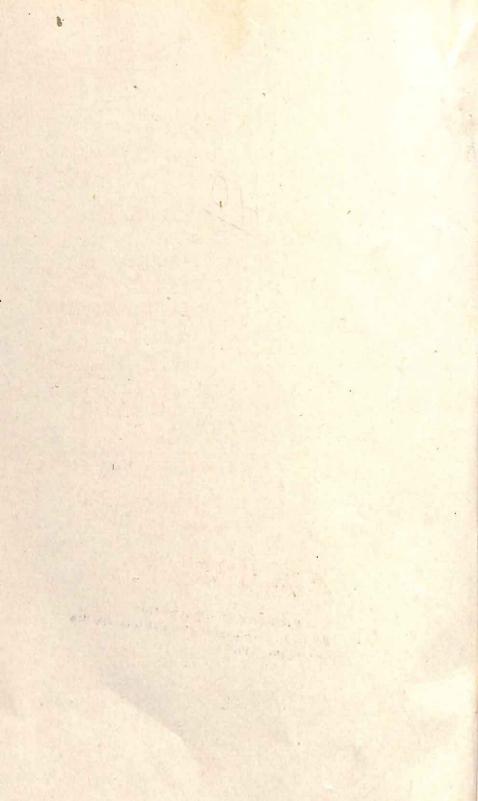
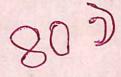


R.K.KAUSHIK M.S.YADAV





DICTIONARY OF CHEMISTRY



by

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UBS Publishers' Distributors Ltd 8/1-B, Chowringhee Lane, Calcutta-700 016 Ph:212451, 249473

1989

ANMOL PUBLICATIONS

NEW DELHI-India

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First Edition: 1987

Reprint : 1989

ISBN 81-7041-041-X

Published by:

ANMOL PUBLICATIONS 4378/4B, Gali Murari Lal, Ansari Road, New Delhi-110002. Ph. 261597.

Distributed by:

Anupama Publishers, Distributors 4378/4B, Gali Murari Lal, Ansari Road, New Delhi-110002. Ph. 261597.



Uns redustions' Distribution ted 847-0, Chowelephop Lane, Calcutta- 200 016 PN 2322381, 249473

Printed at :

MEHRA OFFSET PRESS, NEW DELHI.

Preface

In this dictionary, an attempt has been made to provide an explanation of the terms used in the various branches of chemistry, together with brief accounts of important substances and chemical operations. We have necessarily had to be selective rather than comprehensive.

An attempt has been made to write the entries in a clear and lucid style to provide both straight forward definitions and invaluable background informations. Structures of complex compounds have been included. The network of cross-references has been maintained throughout this book. The emphasis in this dictionary is placed on providing definitions rather than on pronounciation, etymology or syllabication. In addition to definitions, synonyms, acronyms and abbreviations are given under the appropriate entry.

The dictionary will be of immense value to students of chemistry and to biochemists, scientists and others studying or working in related fields.

The authors are happy to give credit here to all those who had a hand in getting the dictionary into its present form and particularly to put on record our great debt to Dr. G. R. Chhatwal, for the care and interest he took in clearing up complex points so that they might be put in simple English.

In compiling a dictionary of this kind it becomes necessary to draw upon the work of many authorities and seek the advice of colleagues to all of whom the author is deeply indebted.

Finally the authors express their sincere thanks to the publishers and printer for printing this book promptly.

All comments from users on omissions or shortcomings will be most welcome.

September, 1987

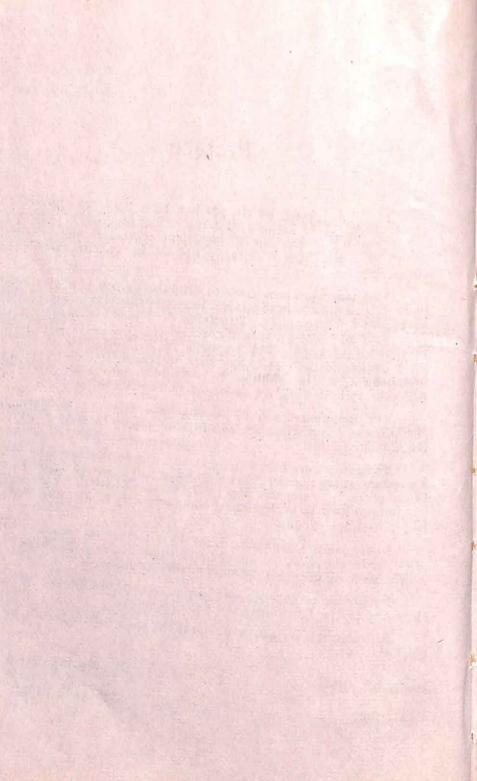
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A

- Abelite. Explosive made up of ammonium nitrate and trinitrotoluene; some forms contain starch and sodium chloride or dinitrobenzene and sodium chloride.
- Abietic Acid. C₂₀H₃₀O₂. M.P. 158°, an -68 to 69°. A crystalline tricyclic triterpene derivative, prepared from plant resins. A monocarboxylic acid and forms retene or 1-methyl-7-isopropyl-phenanthrene on dehydrogenation. A chief constituent of colophony. It is present in various varnishes. It is used in detergents.
- Abrin. A toxic substance. It causes agglutination of the red blood corpuscles and present in the seeds of the Abrus precatorius, an Indian shrub. It is a mixture of proteins.
- ABS Detergent. (Sodium Alkylbenzene Sulphonates). Manufactured by a Friedel-Craft alkylation of benzene with long chain alkenes $(C_{10}-C_{12})$. It is used in powder detergents (e.g., Surf).
- Absolute Alcohol (ethanol). Pure alcohol, C₂H₅OH, a colourless liquid with a pleasant odour, B.P. 78 32°, Density 0.7936 at 15°. Miscible with water. It acts as a solvent for many organic compounds. It is obtained by reducing acetaldehyde by sodium amalgam in water. It is used in the preparation of ether and chloroform. Detected by oxidation to acetaldehyde with potassium dichromate.
- Absolute Temperature. A temperature on the "absolute" (or Kelvin) scale; symbol TA; the zero of this scale is the temperature at which a perfect gas would have zero volume if it could be cooled indefinitely without liquefaction or solidification. Temperature is defined by the relationship:

$T = \theta + 273.15$

where θ is the celcius temperature.

- Absolute Zero. The zero value of thermodynamic temperature; C kelvin or -273.15°C.
- Absolute Configuration. It represents the arrangement of atoms or groups about the carbon atom.
- Absorbance (Molar Extinction Coefficient). It is the efficiency of different organic structures to trap a photon and proceed to the excited state; expressed a € and increases with increasing conjugation in a polyene.
- Absorptiometer. An apparatus for determining the solubility of a gas in a liquid.

- Absorption. A process in which a gas or vapour is taken up by solid or liquid or in which a liquid is taken up by a solid. The absorbed substance distributes itself in the bulk of the material. Selective absorption is used to separate components of mixtures of gases or vapours.
- Absorption Bands. The bands observed in an absorption spectrum are known as absorption bands.
- Absorption Co-efficient of a Gas. It is the volume of a gas measured at 0° and 760 mm pressure, which will dissolve in 1 cc. of a liquid.
- Absorption Co-efficient of Light. See Lambert's and Beer's Law.
- Absorption Indicator. An indicator used for titrations that involve a precipitation reaction. Fluorescein—a fluorescent compound—is commonly used as absorption indicator in sodium chloride and silver nitrate titrations. Sodium ions and chloride ions are fluorescein (negative ions) ions are absorbed by silver chloride precipitate forming a pink complex.
- Absorption of Light. When the light beams fall on the surface of a transparent substance, part of the light is reflected and the rest is transmitted unchanged. However, the light is neither reflected nor transmitted but absorbed when falls on an opaque surface. Since, this surface absorbs light of all wavelengths, the process is called absorption. If a light of particular wavelength is used, with the return of the atom or molecule from a state of lower inducing chemical reactions.
- Absorption Spectrum. Absorption spectra are obtained by passing white light through the atomic/molecular vapour, which absorb radiations of the characteristic frequencies, thereby, producing dark lines on a white background. The position of the lines in exactly the same.
- Absorption Tower. It is a part of the plant where the sprayed liquid dissolves the gas being passed counter-currently up the or coke or Raschig rings. These are also known as Scrubbers.
- Abundance. The relative amount of a given element amongst 50% by mass.
- Abyssinian Gold. It is a yellow alloy of 90.7% Cu and 8.3% Zn plated with gold on one side and rolled into sheets.
- Accelerator. A catalyst added to reactants to increase the rate of

Acceptor. The atom or group to which a pair of electron is donated in a co-ordinate bond. For example, the formation of ammonium ions from ammonia can be represents as:

$$H: N: + H^{+} = \begin{bmatrix} H & \cdots & H \\ H: N: H \end{bmatrix}^{+}$$

$$H$$
Donar Acceptor

Accumulator (Secondary cell, Storage battery). It is a reversible electric cell or battery that can be charged by passing an electric current through it. A common example is the lead acid accumulator, used in vehicle batteries.

Acenaphthene.
$$C_{12}H_{10}$$
 $\left(\begin{array}{c} CH_2-CH_2 \\ \end{array}\right)$. Colourless needles, M.P.

95°, B.P. 278°, soluble in hot alcohol. On oxidation with acid dichromate forms naphthalic acid and when passed through a red hot tube forms naphthylene. Forms picrate, M.P. 161°. Obtained from coal tar and synthesised from alcoholic potash and α-bromoethylnaphthalene. It is used as an intermediate in dyestuffs.

sparingly soluble in water. It produces dyes like; Ciba Scarlet G, Ciba Red R, etc.

Acenaphthylene,
$$C_{12}H_8$$
 (Yellow plates, M.P. 92°-

93°, B.P. 265°-275°. Insoluble in water but soluble in alcohol and ether. D15 0.899. Recrystallised from alcohol.

Acetal, CH₃.CH(OC₂H₅)₂. A form of organic compound obtained by an addition of an alcohol to an aldehyde. A pleasant smelling liquid, B.P. 104°=105°, soluble in alcohol and ether, partially miscible with water. It can be prepared by passing acetylene into alcohol in the presence of a catalyst a by mixing aldehyde and ethanol.

- Acetaldehyde, CH, CHO. A colourless liquid with a characteristic smell. B.P. 20.8°, D. 0.801 soluble in water, alcohol and ether; insoluble in concentrated calcium chlorode. Prepared by oxidation of alcohol with K2Cr2O7 and sulphuric acid. Manufactured by passing alcohol and air vapours over heated Cu or Ag. It forms Aldol. It is used for the manufacture of
- Acetaldoxime, CH₃CH=N-OH. White crystalline needles M.P. 47°, B.P. 115°. Soluble in water, alcohol and ether. Prepared from hydroxylamine and aqueous solution of acetaldehyde. It is used
- Acetamide, CH₃CONH₂. All white needles, absorbs water and liquefy, strong odour of mice, M.P. 82°, B.P. 222° soluble in water and alcohol. Prepared by the dry distillation of ammonium acetate. Forms amides (salt like compounds) with
- Acetanilide, C₈H₉ON. White crystals, M.P. 114°, B.P. 314° soluble in alcohol, ether and chloroform. Prepared by refluxing excess acetic acid with aniline. It is known as an antifebrin, antipyretic
- Acetanin, C₁₄H₈O₇ (O.COCH₃)₂. A greenish white soluble in ethylacetate. Prepared from tannic acid and acetic anhydride. Used in chronic diarrhoea and intestinal catarrh.
- Acetarsol, (3-acetalyl amino-4-hydroxy-phenyl arsonic acid): CH₃.CO.NH.C₆H₃(OH)AS.O(OH)₂. White crystals, M.P. 240° – 250°. Soluble in alkalis. Prepared from 3-nitro-4-hydroxyphenylar sonic acid, used in dysentery, malaria and diseases due to
- Acetates: Salts or esters of acetic acid obtained by the action of
- Acetate Silk Dyes. These dyes are ester like compounds soluble in organic solvents. These are used for dyeing acetate silk. (cellulose acetate). Three types of dyestuff:
- 1. Ionamines. The formaldehyde bisulphite compounds of some 2. Dispersols.
- These are fat-soluble dyes and are used in colloidal solution.
- 3. Duranols. Aminoanthraquinones used in colloidal solution.
- Acetic Acid, Glacial Acetic Acid, CH3COOH. A colourless liquid with pungent smell, M.P. 16.7°, B.P. 118.5°, D²⁰ 1.0491. Soluble in water, alcohol and ether. Manufactured by oxidising actialdehyde by air at 60° and high pressure in presence of manganese acetate. It does not attack Sn. Forms acidic, basic and normal salts with metals. Nitric acid and chromic acid

do not oxidise it. Used in the preparation of metallic acetates and esters; in the manufacture of cellulose acetate and as an ine t solvent for many organic compounds.

Acetic Anhydride, C₄H₆O₃. A colourless liquid with a pungent smell, D²⁰ 1 081, B.P. 139 5°: 12% soluble in cold water. Miscible with ether and benzene. Manufactured by passing acetylene into acetic acid in presence of Hg salts or by passing the acid vapours over a heated catalyst, or more easily by heating the vapours to 600°—1000°. It forms acetyl compounds with compounds containing -OH and -NH₂ groups. Used in the manufacture of aspirin and cellulose.

Acetins. They are acetates of glycerol. There are three main types of acetins:

- Monoacetin, CH₂OH.CHOH.CH₂OOC.CH₃.D²⁰ 1'221, B.P. 130° -132° at 2-3 mm. It is quite soluble in water and chloroform. Prepared by heating glycerol and acetic acid with sulphuric acid; used as a solvent for dyes used in printing paper bags.
- 2. Diacetin, CH₃COOCH₂CHOH.CH₂OOC.CH₃.D²⁰1·16, B.P. 259°. Soluble in water and alcohol but insoluble in ether and water in presence of water. Formed as a by-product in the formation of triacetin. Used as a solvent for basic dyes and plasticizer for cellulose acetate lacquers.
- 3. Triacetin, D²⁰ 1'16—1'17, B.P. 260°. Soluble in ether. Obtained by heating glycerol with excess of acetic acid and extracting the triacetin from aqueous solution with ether. Used as a solvent for gums and resions and as a plasticizer for lacquers.
- Acetoacetic Acid, Acetone Carboxylic Acid, C₄H₆O₃, CH₃CO. CH₂CO.OH. A colourless syrup, decomposes into acetone and CO₂ below 100°. Obtained from acetoacetic ester. It is present in abundance in the urine of diabetic patients. Aqueous solution of the acid forms a violet colour with ferric chloride.
- Aceto Acetic Ester, Ethyl Acetoacetate, CH₃COCH₂COOC₂H₅. A colourless mobile liquid with a pleasant smell, D²⁰ 1'0256, 181°-182°. Miscible with alcohol, ether and benzene. It is prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium or sodium ethoxide on acetic prepared by the action of sodium ethoxide on acetic

- Acetoin, Acetyl Methyl Carbinol, C₄H₈O₂.CH₃CH(OH)COCH₃. A colourless liquid, M.P. 15°, B.P. 148°. Miscible with water and alcohol and immiscible with ether. Obtained by reducing diacetyl. Reduces Fehling's solution and forms crystalline compound with sodium bisulphite.
- Acetol, Hydroxyacetone, Pyroracemic Alcohol, Pyruvic Alcohol, C₃H₆O₂, CH₃CO.CH₂OH. A colourless liquid with a pleasant odour, D²⁰ 1 082, B.P. 145°-146°. Miscible with water, alcohol and ether. Obtained by the action of Bacterium Xylinum on a-propylene glycol. Methyl alcohol prevents its polymerisation. It reduces ammoniacal silver nitrate in cold. Detected by fluorescence when reacted with o-aminobenzaldehyde.
- It is a process of removing acetyl group from an Acetolysis. organic compound by heating it with aqueous or alcoholic
- Acetone, Dimethyl Ketone, CH₃COCH₃. A colourless volatile liquid with a pleasant smell, highly inflammable, D20 0.8186, B.P. 56°. Soluble in water, alcohol and ether. Sparingly soluble in calcium chloride solution. Manufactured from propene, either by the air-oxidation of propan-2-ol or as a by-product from cumene process. Used as a solvent and in the manufacture of 2-methyl-propanote. Detected by the formation of lodoform, a red colour with sodium nitroprusside and ammonia which changes to violet on addition of acetic acid. It is an excellent solvent for acetylene, and is used in cylinders of the
- Acetone—Chloroform, Chloretone, (CH₃)₂C.OH.CCl₃. A white camphor like smell. M.P. 97°, B.P. 167°, C.H. and acetone obtained by reacting KOH and acetone and chloroform mixture in cold. It is steam
- Acetone Dicarboxylic Acid, β-ketoglutaric Acid, CO.(CH₂COOH)₂. Colourless needles, M.P. 135, soluble in water and alcohol, insoluble in C₆H₆ and CHCl₃, obtained by the action of sulphuric acid on citric acid. Forms violet colour with FeCl₃ and a white ppt. with mercuric sulphate. It is used in organic
- Acetonyl. CH₃COCH₂—radical is known as acetonyl.
- Acetonylacetone, 2:5 Hexadione, CH3COCH2CH2COCH3. A colourless liquid turning valley. colourless liquid turning yellow on standing. D20 0.970, B.P. 191.3, soluble in water, alcohol and ether. Obtained as a by-product in the manufacture of acetic acid from acetylene-It is used as a solvent for cellulose acetate.

Acetophenone, C₈H₈O, C₆H₅COCH₃. Colourless plates, M.P. 20°C, odour of bitter almonds. Prepared by the action of ethanoyl chloride upon benzene in the presence of AlCl₃. Shows characteristic properties of ketones. Oxidised by potassium permanganate to phenylglyoxalic acid. Forms oxime, M.P. 60°C, and phenylhydrazone, M.P. 105°C. Used as a solvent for cellulose ethers and as a soporfic (hypnone).

Acetoxy. The group CH₃COO-.

Acetylacetone. Metal derivatives of acetylacetone usually containing the grouping shown (Macac) with some delocalization in the ring. The number of acetylacetonate groups bonded to the metal can vary from 1 to 4.

Acetylation. The process for introducing acetyl groups into an organic compound containing -OH or -NH₂ or -SH groups. It is carried out by heating the compound with acetic anhydride (ethanoic anhydride) or ethanoic chloride usually in presence of an inert solvent like benzene or acetic acid. In many cases zinc chloride or benzene is used to increase the rate of reaction.

Acetyl Chloride, CH₃COCl. A colourless liquid with pungent odour, fumes in moist air, producing acetic and hydrochloric acids. D° 1·130 C, B.P. 55°C. Soluble in ether and benzene, reacts with water and other —OH compounds. Prepared by the distillation of a mixture of acetic acid and phosphorus trichloride or oxychloride. Used to prepare acetyl derivatives of compounds containing —OH and —NH₂ groups.

Acetylcholine, C₇H₁₇NO₂, (CH₃)₃N⁺CH₂CH₂OOC CH₃OH⁻. A white hygroscopic crystalline powder, soluble in water. Occurs in blood, ergot and in some plants. If injected, causes dialation of arteries. Passage of a nerve impulse from a nerve end to another nerve or muscle cell liberates acetylcholine. Causes a rapid fall in blood pressure, quickly destroyed by the hydrolytic action of acetylcholine esterase.

Acetylcoenzyme. An important thioester of importance in metabolism and biosynthesis. Acetyl co-enzyne A is the main substrate of the citric acid cycle. Acts as an acetylating agent in the synthesis of acetylcholine. Used in the biosynthesis of: (1) fatty acids (2) polyketides and (3) terpenoids and steroids. Acetylene, C2H2, CH=CH. A colourless gas with a pleasant smell, M.P. -82°C, B.P. -84°C, liquified below 37°C and solidified in liquid air. Soluble in several organic solvents and in water; acetone absorbs about 25 times its volume at 15°C and 760 mm. Both gas and liquid are very explosive. Stored and transported in steel cylinders containing propanone and a porous material such as diatomaceous earth. Prepared, generally, by reacting calcium carbide and water or by the pyrolysis of the lower paraffins in the presence of steam or by the partial oxidation of natural gas (methane) and by the cracking of hydrocarbons in an electric arc. Used for the preparation of acrylonitrile, vinylchloride, vinyl acetate, ethanol, ethanoic acid, neoprene and polyvinylalcohol. Also used in oxyacetylene weldings.

Acetylene Black, Cuprene. A form of carbon black obtained by pyrolysis of ethyne (acetylene).

Acetylene Dichloride. See dichloroethenes.

Acetylene Tetrachloride, Sym-Tetrachloroethane, HCCl2.CHCl2. A colourless liquid with chloroform like smell, D20 1.60, B.P. 146'3°C, insoluble in water but miscible with benzene, volatile in steam. Manufactured by passing chlorine separately into a solution of antimony pentachloride in acetylene tetrachloride (tetrachloroethane). Reacts with dilute alkalis to

Acetylides. Carbides containing C_2^{2-} or C_2R^- species. Formed by more electropositive elements (e.g. K, Ca, Al) and by several transition elements (e.g. Cu, Ag, Au). Most transition metal acetylides are explosive. Hydrolysed to acetylene. Form complex acetylides (e.g. $[M(C \equiv CR)_n]^{\times -}$) with transition metals

Achiral. The molecule of a compound which is optically inactive.

Achromatic Indicators. Substances used in the titrations of turbid

Acid. An acid produces H+ ions when in aqueous system. Most inorganic acids are compounds of an acidic oxide and water. Some acids do not contain oxygen (e.g. HCl, HF, H2SiF6 etc.) while some acids contain oxygen (e.g. HNO₃, H₂SO₄ etc.). In inorganic acids, the oxide concerned (of a metal) may exhibit amphoteric character. Aqueous solutions of acids show sharp task, turn blue litmus red, liberate CO2 from a metallic carbonate and show reactions characteristic of the anion present. Aqueous solution of an acid contains the hydroxonium ion, H₃O+ (as proton can not exist in free state). Acids can also exist in non-aqueous solvents. NH4+ ion is formed by the solvation of a proton by ammonia. Substances which dissolve in ammonia to form the ammonium ion are acids in the behaviour e.g.

NH₄Cl. Liquid water ionizes to form hydroxonium and hydroxyl ions:

 $2H_2O \rightleftharpoons H_3O^+ + OH^-$

This ionization being the reverse of the ionization reaction in water, substances producing hydroxyl ions in water are bases. Liquid ammonia ionizes:

 $2NH_3 \rightleftharpoons NH_4^+ + NH_2^-$

and ionic amides so formed are bases in the process. The solvents which do not contain hydrogen but produce an appropriate positive ion is an acid in the process. Bromine trifluoride ionizes:

 $2BrF_3 \rightleftharpoons BrF_2^+ + BrF_4^-$

and a substance forming the BrF₂⁺ ion in solution e.g. BrF₃. SbF₅, is an acid in the system.

Typical organic acids contain the group —COOH (e.g. CH_3COOH , C_6H_5COOH etc.) but many other organic compounds [containing groupings e.g. the sulphonic —S(O)₂.OH show acidic properties. Carbolic acid, C_6H_5OH , and picric acid, $C_6H_2(NO_2)_3OH$, behave like acids, but are not considered acids.

According to Lewis, acids are electron acceptors. Thus AlCl₃ can accept electrons from a chloride ion producing the (AlCl₄) ion and is a Lewis acid. The degree of ionization of an acid is a measure of its strength. Strong acids are fully ionized while weak acids are feebly ionized.

Acid Dyes. Dyes have an aromatic chromophoric group and a group soluble in water (e.g. SO₃H group as its sodium salt). The acid dyes are of three types.

Simple acid dyes. contain no polyvalent metals and are not improved by after treatment with Na₂Cr₂O₇/HCl (dil).

Mordant acid dyes, generally, contain ortho OH—azo or OH—OH groups and combine with the mordants [e.g. Cr(OH)₃] and the fibre.

Premetallized acid dyes are as simple as acid dyes.

Acid dyes are good for wool but not for cotton.

Acid Egg. Device, made of acid-resistant material, into which acid is run, to be distributed by compressed air.

Acid Exchange Resins. Synthetic resins, which have the property of absorbing anions from material in contact. Used for removing impurities and unwanted salts from aqueous solutions.

Acid Oil, Cresylic Acids or Phenols. The alkali extract of phenol derivatives obtained during the formation of gasoline by cracking method.

Aconitic Acid, C6H6O6. Unsaturated, colourless leaflets, M.P. 194°C. soluble in water, prepared by dehydrating citric acid with 50% sulphuric acid.

Aconitine, C₃₀H₃₅O₅(OCH₃)₄(OH)₂N. Colourless hexagon crystals, M.P. 204°C, slightly soluble in water, $(\alpha)p+17\cdot3^{\circ}$ in CHCl₃. Highly poisonous alkaloid obtained from Aconoitum napellus.

Colourless needles

M.P. 111°C, soluble in alcohol.

Acriflavine, 2:8 diaminoacridine methochloride, C14H14N3Cl. Orange red crystals, soluble in water, prepared by methylating proflavine, a mixture of 2, 8-diaminoacridine dihydrochloride and 2, 8-diamino-10-methylacridinium chloride hydrochloride, used for wound dressings.

Acrilan. A brand name for a synthetic fibre.

Acrolein, Acraldehyde, C3H4O, CH2=CH-CHO. Colourless volatile liquid, poisonous vapours, D²⁰ 0.841, B.P. 52°C, soluble in water and alcohol. Prepared by distilling a mixture of glycerine, potassium sulphate and potassium bisulphate.

Acredein Polymers, Propenal Polymers. Polymers of CH2=CH. CHO, formed by free radical polymerisation. Used as thickening agents, protective colloids and in plastics and lacquers. Disacryl is insoluble in acrolein polymer.

Acrylate Resins and Plastics. Referred to polymeric methyl acrylate (methylpropenoate) and polymeric methyl methylacrylate (methacrylate) (methyl 2-methyl propionate), used in emulsion form in textile and leather finishes, lacquers paints, adhesives and safety glass interlayers. Polymeric methyl methyl acrylate forms Perspex.

Acrylic Acid, Vinyl Formic Acid, C₃H₄O₂, CH₂=CH-COOH. A colourless liquid, D¹⁶ 1.002, M.P. 13°C, B.P. 141°C, miscible with water. Prepared by oxidising acrolein with moist silver

Actinides. The elements actinium, thorium, protactinium, uranium neptunium, plutonium, americium, curium, berkelium, californium, einsteinium, fermium, mendelevium, nobelium, and lawrencium collectively are termed as actinides, 5f shell in filled and are analogous to lanthanides, radioactive in nature and handled by remote control, all elements above atomic number 93 are artiActinium, Ac. At. no. 89, M.P. 1050° C, B.P. $3200\pm300^{\circ}$ C, occurs in uranium ores, prepared by bombardment of radium with neutrons, the isotope ²²⁷Ac is very radioactive ($t_{1/2}$ 22 years), the metal glows blue, also prepared by reducing AcF₃ with Li vapours at 1200° C, silvery white in colour, forms tripositive compounds.

Activated Carbon. See active carbon.

- Activation Energy. Molecules possess average energy which is insufficient to enable them to undergo chemical reactions. The excess energy, above this energy, required by the molecules for reaction to proceed is known as activation energy (E_a).
- Activator. A substance added to accelerator enhances its activity to full activity e.g. ZnO.
- Active Carbon. Carbon (charcoal) treated at a high temperature with steam air or CO₂ (activation of carbon). Used as an adsorbent (due to large surface area and opened capillaries) for removing small traces of impurities from a gas or liquid, for water waste and water treatment, air pollution control, sugar refining, dry cleaning, rubber reclamation, cigarette filters and as a catalyst.
- Active Centres. The specific sites on a catalyst surface at which absorption occurs.
- Active Mass. The thermodynamic activity and is a product of the affinity of a reacting substance and its concentration. Consider the reversible reaction and the extent to which the reaction

A+B=C+D

proceeds is proportional not to the affinities, but to the active masses of the reacting substances and the equilibrium constant K is given by:

 $K = \frac{[C][D]}{[A][B]}$, where [C][D][A] and [B]

are the active concentrations of products and reactants respectively.

- Activity. A thermodynamic quantity and a measure of the effective concentration of a particular substance in a given chemical system. Two types of activity are:
 - (i) Absolute activity, a° is given by $\mu = RT \ln a^{\circ}$, where μ is the chemical potential of the substance, R is the gas constant, T is the absolute temperature.
 - (ii) Relative activity, a, is given by $\mu = \mu^{\circ} + RT \ln a$, where μ° is the chemical potential of the substance in its standard state.

Activity is proportional to partial pressure of the gas for ideal gases and directly proportional to the concentration for dilute and ideal solutions.

- Act ivity Coefficient (f or γ). Thermodynamic activity, (a) in a system is given by a=fc or $a=\gamma c$ where c is the concentration of the substance. For ideal or dilute solutions, for $\gamma = 1$. For electrolytes, activity co-efficients are considered to be the geometrical mean of the single ion activities.
- Activity Series. A series in which the elements are arranged in order of their electrode potential.
- Actomyosin. An important protein of body muscle.
- Acyclic. Organic compounds containing chains without rings but possibly with branches.
- Acyl. A general name for organic acid radicals, which are the residues of carboxylic acids after removal of the -OH group e.g., acetyl chloride, CH₃COCl obtained from CH₃COOH. The names of the individual acyl radicals are given by replacing the -icof the acid by -vl.
- Acylation. A process which substitutes the acyl (RCO-) group into a molecule, generally for an active hydrogen of e.g. an -OH group.
- Acyloins. α-β-Ketoacids of the form R.CO.CHR'.OH, prepared by acyloin condensation, a process of condensing two molecules of
- Adamantane, C10H16. Colourless hydrocarbon, M.P. 269°C, sublimes at room temperature and pressure, has a rigid ring system constituted of three fused cyclohexane rings, configuration similar to diamond lattice, synthesised from tetrahydrodicyclopentadiene. 1-Adamantane hydrochloride is useful in viral infections and Parkinson's disease. Its derivatives are used as lubricants
- Adatom. An adsorbed atom.
- Addition Reaction. A reaction leading to the saturation or part saturation of an unsaturated system (C=C, C=C, C=O, etc.) by addition of a molecule across the multiple bond. For example, formation of 1, 2-dibromoethane from bromine and ethane, the Diel's-Alder reaction and addition polymerisation. Also used in inorganic chemistry e.g., the reaction of BF3 with nucleophiles, such as, ammonia, ether to form complexes.
- Additive Volumes, Law of. The volume occupied by a mixture of gases is equal to the sum of volumes which would be occupied by the components under the same conditions of temperature and pressure.
- Adduct. A phase (generally a compound) obtained by direct combination of two or more different compounds or elements.
- Adenase. The enzyme responsible for deaminating adenine to hynox-

- Adenine, 6-Aminopurine, C₆H₅N₅. Long rhombic needles with three molecules of water which it loses at 110°C. A constituent of the nucleic acid portion of nucleoproteins, and, combined as adenyl-pyrophosphate. M.P. 360°-365°C, plays an important part in many metabolic processes.
- Adenosin, Adenine Riboside. Long thin needles, M.P. 229°C, soluble in water, insoluble in alcohol, (α) p—63·3°. A nucleoside made up of one molecule of adenine and one molecule of pentose D-ribose, forms a part of the structure of ribose nucleic acid.
- Adenosine Diphosphate (ADP). Adenosine 5'-diphosphate and a precursor of ATP, also obtained from it during process in which ATP is involved.
- Adenosine Monophosphate (AMP). Generally, adenosine 5'-phosphate (muscle adenylic acid), an important structural component of nucleic acids and of various co-enzymes.
- Adenosine Triphosphate (ATP). The most important of the so called 'high energy compounds', a group of naturally occurring organic phosphates having high free energies of hydrolysis, and playing a basic part in biosynthesis, active transport and muscle action. A primary source of energy in the metabolism of plant, animals and bacterial cells.
- Adenylic Acid, Aldenosine Phosphoric Acid. A nucleotide made up of a molecule of adenine and phosphoric acid connected through D-ribose.
- Adhesion Agent. An additive improving the adhesivity of a material, prevent stripping of an adhesive from a surface by water, used in road bitumens, surface dressings and in coated macadam surfacings.
- Adhesive. A liquid material which joins two surfaces by forming a solid join.
- Adiabatic Change. A change in which no heat transfer takes place in a system.
- Adipic Acid, C₆H₁₀O₄, HOOC. [CH₂]₄. COOH. M.P. 153°C, present in beet-juice, prepared by the oxidation of cyclohexane by air, forms long chain polymers (e.g., Nylon), used to manufacture plasticizers and some vinyl and urethane plastics.

Adrenaline, Epinephrine, C₉H₁₃NO₃. CH.OH.CH₂NH.CH₃

M.P. 212°C, synthesised from catechol. Used as the acid tartarate in the treatment of allergic reactions, and circulatory collapse, local anaesthetic injections.

Adsorbate. A substance adsorbed by the surface of absorbent.

Adsorbent. A substance whose surface adsorbs the absorbate.

Adsorption. A process in which a layer of atoms or molecules of one substance forms on the surface of a solid or liquid. The adsorbed layer may be held by chemical bonds (chemisorption) or by weaker van der Waals forces (physiosorption). Adsorption process may involve either simple unimolecular adsorbate layer or multilayers. The process is useful in processes such as the purification of materials, drying of gases, control of factory effluents, production of high vacua, etc. Adsorption processes are basis of colloidal and emulsification behaviour and heterogeneous catalysis.

Adsorption Indicator. An indicator which works by adsorption on the surface of a precipitate, e.g., fluorescein in the volumetric estimation of Ag⁺ ions by Cl⁻ ions.

Aerosol, Aerogel. A dispersion in which a finally divided solid is suspended in air and the size of particles are of colloidal dimension e.g., smoke, used in hair lacquers, paints etc. It is also a trade name for wetting agents such as esters of sodium sulphosuccinic acid.

Aetiophyllin. See chlorophyll.

Affatoxins. A toxic metabolites of the fungus.

Aspergillus Flavus. Link ex-Fries which is present in grain crops and peanuts, extremely toxic and

carcinogenic, exists in four isomers [B1, B2, G1 and G2].

Ag. Silver metal.

Agar-agar. A seaweed colloid. A mixture of agarose and agarospectin, extracted by hot water from various marine algae Rhodophycea in the powder form, dissolves in hot water and set to a jelly (0.5% concentration), used as a medium for cultivating bacteria and moulds, as a laxative, as a thickner and as an emulsion stabilizer in the foods.

Agate. A variety of SiO₂ with purple or brown coloured bands, used as a gemstone for making mortars and pestles, and as a bearing surface in scientific instrument.

- Aggregate. Pieces of material which are, or can be, united by a cementitious substance or bond to form a solid mass. In concrete the aggregate consists of crushed stone, slag or clinker and sand, united by portland and other cements. In bricks and other ceramic materials, particles of non-plastic materials form the aggregate; they are united by plastic clay when in the green or unfired state and later (in the burned state) by a crude glass which fuses and on cooking, solidifies and form bonds. Properties of the product are influenced by the size and shape of the product.
- Agiucone, Aglycone. The term collectively used for the non-sugar portion of a glycoside which breaks off from the sugar on its hydrolysis by acid or enzyme.
- Air. The volume composition of normal dry air is N_2 (78.08%), O_2 (20.94%), Ar(0.934%), $CO_2(0.03\%)$, Ne(0.0018%), Kr(0.0001%), Xe (0.000009%), Ru (6×10⁻¹⁸%). The physical constants of dry air are as follows: D 1.2928 gl⁻¹ at 273 K and 760 mm; Cp 0.2396; Cv 0.1707; R 2.1529×10³ (pressure mm; Cm^3 g⁻¹). Air can be liquified by rapid expansion into a partial vacuum (pale blue liquid), B.P. $-192^{\circ}C$ (air) to $-185^{\circ}C$ (after N_2 evaporation), D 0.9 (approximately).
- Air Filters. Filters constructed to remove atmospheric dust from an air stream, generally, they are composed of a space packed with a filting medium, such as glass fibre or slag wool through which air is passed.
- Air Hardening. A method of hardening steel by allowing it to cool in a natural way in air (or in an air blast), also known as air quenching.

Air Lift Pump. An apparatus for pumping corrosive liquids.

Al. Aluminium metal.

α-Alanine, 2-Aminopropanoic Acid, C₃H₇NO₂, CH₃CH(NH₂) COOH. M.P. 200°C, rhombic prisms, soluble in 4—5 parts of water, insoluble in alcohol, [α_D]²²+2.58°, one of the aminoacids obtained from the hydrolysis of proteins.

β-alanine, β-aminopropionic acid, C₃H₇NO₂, H₂N. CH₂. COOH. M.P. 196°C, soluble in water, prepared by the action of sodium hypobromite on succinimide, used in the preparation of pantothenic acid (vitamin)

Albumins. A class of proteins, soluble in water and coagulated by heating e.g., ovalbumin (from eggs) and lactalbumin (from milk).

Alcian Blue. A dyestuff, Alcian Blue 8 G.S.

Alcohol. Organic compounds of the general formula ROH, where R is a hydrocarbon group e.g., methanol (CH₃OH) and ethanol (C₂H₅OH). Three classes of alcohols are:

primary alcohol (carbon atom attached to two hydrogen

secondary alcohol (carbon atom is attached to one hydrogen atom

and two other groups),
$$-C-C-C-e.g.$$
, $CH_3.CH(OH).CH_3$.

tertiary alcohol (carbon atom is attached to three other;

groups,
$$-C-C-C-\hat{e}.\hat{g}., CH_3.C(OH)-CH_3$$

 $-C CH_3$

Alcohols can be prepared by:

1. Hydrolysis of haloalkanes using aq. KOH:

$$RI + OH^- \rightarrow ROH + I^-$$

Reduction of aldehydes by nascent hydrogen (from Na/Hg):
 RCHO + 2(H) → RCH₂OH.

The important reactions are:

1. Oxidation

2. Ester formation

3. dehydration (over hot munic store)

$$\begin{array}{c} R CH_2CH_2OH \xrightarrow{-H_2O} R CH = CH_2 \\ alkene \end{array}$$

There are monohydric alcohols, containing one—OH group (e.g., CH₃OH), dihydric alcohols, containing two—OH group (e.g., CH₂—OH

CH₂—OH, glycol) and trihydric alcohols, containing three —OH.

- Aromatic derivatives having -OH group attached directly to carbon atoms of the ring are called *phenols* and show distinct property (e.g., C_6H_5OH , phenol).
- Aldehyde Polymers. Polymers with a CHRO backbone formed by polymerization of aidehydes RCHO (e.g., polyformaldehyde and polychloral). Used in machine parts, car parts, etc.
- Aldehydes. A type of organic compound with the general formula RCHO, where the —CHO group (the aldehyde group) consists of a carbonyl group attached to a hydrogen atom, e.g., HCHO (formaldehyde), CH₃CHO (acetaldehyde). Colourless liquids (aliphatic) or solids (higher aromatic) with characteristic smell, formed by oxidising a p-alcohol, oxidised to acids and reduced to p-alcohols. Aromatic aldehydes are prepared by heating aromatic hydrocarbons with CO, HCl, anhydrous AlCl₃ (Gattermann-Koch Synthesis). Alkaline solutions of phenol react with CHCl₃ (Reimer-Tiemann reaction) to form phenotic aldehydes.

Aliphatic and aromatic aldehydes form adducts with alkali bisulphites, and form cynohydrins, aldoximes, semicarbazons and phenyl hydrazones, form acetals with alcohls, undergo aldol condensation, form Schiff's bases with p-amines, they reduced Fehling's solution and Tollen's reagent, some of the aliphatic and aromatic aldehydes undergo Cannizzaro reaction.

- Aldicarbs, C₇H₁₄N₂O₂S. An insectiside and nematocide commonly used on Sugarbeet and potatoes.
- Aldol, acetaldol, 3-Hydroxybutnal, C₄H₈C₂, CH₃.CH OH.CH₂CHO A colourless oily liquid, B.P. 83°C/20 mm, manufactured by treating ethanol (C₂H₅OH) with alkali carbonates, lime, borate, etc., reduction under pressure forms 1, 3-butylene glycol.
- Aldol Condensation. A reaction between two molecules of aldehyde (aliphatic) to form a β-hydroxyaldehyde, e.g., aldol (CH₃CHOH CH₂ CHO) obtained from ethanol in presence of a catalyst (KCN, sodium ethanoate or dil. NaOH), only aldehydes containing the -CH₂.CHO grouping can produce aldols, other aldehydes undergo Cannizzaro reaction.
- Aldomic Acid. An acid derived from an aldose by oxidation of the aldehyde group to a carboxyl group, e.g., gluconic acid.
- Aldose. A sugar containing a potential aldehyde (-CHO) group. The group may be obscured by its inclusion in a ring system. Aldoses are named aldopentoses, aldohexoses, etc., depending on the number of carbon atoms.
- Aldosterone, $C_{21}H_{28}O_5$. The most active steroid hormone secreted by the adrenal cortex.

Aldoximes. Compounds obtained by the action of hydroxylamine on aldehydes, the group present is -CH=N.OH, prepared by the action of p-amines with permonosulphuric acid, aliphatic eximes are colourless liquids or low melting solids, some soluble in water. Aromatic aldoximes are crystalline solids, e.g., benzaldoximes. When boiled with dilute mineral acids, hydroxylamine and the aldehyde is obtained. All aldoximes exist in the

Acid chlorides or anhydrides convert them to nitriles.

Aldrin. The insecticidal product containing not less than 95% by weight of 1, 2, 3, 4, 10, 10-hexachloro-1, 4, 4a, hexahydroexo-1, 4-endo-5, 8-dimethanonaphthalene, obtained from the Diel's-Alder addition product of cyclopentadiene and vinyl chloride by dehydrochlorination followed by condensation with hexachloro cyclo pantadiene, insecticidally active as a contact and stomach poison against a wide range of soil pests, nonphytotoxic but toxic to humans and animals.

Algin. A seaweed colloid, used mainly as the Na salt producing a viscous solution with water. Used as a stabilizer for food products (ice cream, etc.), pharmaceuticals and as a dressing in the textile industry. It is aliginic acid which is D-mannuronic acid.

(where x=200-1000)

Alicyclic. Cyclic carbon eompounds, do not contain any benzene ring, possess, aliphatic characteristics, e.g., cyclopropane, cyclo-

Aliphatic. An organic compound possessing its carbon atoms in chains and not in closed rings, e.g., methane (CH₄), ethanol

Alizarin, C14H8O4. Orange red dyestuff, soluble

in alcohol and ether, M.P. 290°C, forms purple red colour in alkaline solutions from which it is precipitated as red to violet 'lakes' by heavy metal salts. Occurs as a glucoside in madder. produced synthetically by fusing anthraquinone-2-sulphonic acid with NaOH and a little KClO₃. Applied to the mordant fibre, Al(OH)3 forms a bright red lake, Cr(OH)3 a red lake. Fe(OH), a violet lake. Dyeing is always done in boiled water solution.

$$N_2$$
 — OH

m-nitraniline and salicylic acid, used in chromemordanted wool.

Alkali. A hydroxide of one of the "alkali metals" Li, Na, K. Rb and Cs. Even CaO, Ca(OH)2, Na2CO3 give alkaline solutions (pH>7) in water known as a base in aqueous solutions.

Alkaline. An aqueous solution having pH greater than 7.

Alkaline Earth Metals. The elements Mg, Ca, Sr, Ba and Ra which are highly electropositive and produce M2+ ions.

Alkaloids. Organic compounds naturally found in plants, contain oxygen and nitrogen atoms, some of them highly poisonous and produce characteristic physiological actions, vary considerably in chemical properties and constitution, basic in nature and combine with acids to produce salts (water soluble), soluble in organic solvents such as alcohol, CHCl₃ and ether, occur mainly in flowering plants, especially in the Ranunculaceae, Papaveraceae and Solanaceae but generally throughout the botanical sources. Formed from aminoac. Is and methanoate, or methionine, or ethanoate or mevalonate, etc. Many alkaloids are derivatives of pyridine, quinoline or pyrimidine, optically active and dextrorotatory. Examples: morphine, caffeine, nicotine, atropine, cocaine, hyoscyamine, quinine, strychnine.

Alkanals, Aldehydes. Organic compounds possessing terminal R-CHO group.

Alkanes, Paraffins. Aliphatic hydrocarbons of the general formula $C_nH_{2^{n+1}}$ (n is the number of carbon atoms in the molecule), $C_nH_{2^{n+1}}$ (*n* is the number of carbon atoms in the molecule), saturated compounds, first four members are gases, the higher members are liquids and those above $C_{16}H_{34}$ are waxy solids, insoluble in water but soluble in CHCl₃ and C_6H_6 , only Cl₂ and Br, attack, prepared by the following methods:

From a sodium salt of a fatty acid:

RCOO-Na++NaOH → RH+Na2CO2

2. By reduction of a haloalkane:

$$RX + 2[H] \xrightarrow{Zn-Cu/C_2H_5OH} RH + HX.$$

3. By Grignard's reagent :

$$R MgX+H \rightarrow RH$$

The main source of alkanes is natural gas.

- Alkanolamines, Alkyloamines. Hydroxyamines formed by the action of an alkene oxide on aq. ammonia at 50°-60°C, hygroscopic solids with low M.P., usually occur as viscous colourless liquids with ammoniacal smell, used as a catalyst, in rubber.
- Alkenes, Olefins, Unsaturated hydrocarbons with the general formul a C_nH₂ⁿ containing one double bond; isomeric with crycloparaffins. Lower members are gases, the intermediate liquids, higher members are waxy solids. Insoluble in water, soluble in CHCl₃ and C₆H₆, obtained from crude oil by cracking alkanes. Example: ethene (C_2H_4) and propene (C_3H_6) , etc. Reduced to alkanes by hydrogen in presence of catalyst (RCH =CH₂+H₂→RCH₂CH₃), React with halogens to give dihalides $(RCH=CH_2+Br_2\rightarrow RCHBr.CH_2Br)$, oxidation by cold permanganate solution forms diols $(RCH=CH_2+H_2O+O\rightarrow$ RCH (OH) CH₂OH), oxidised to cyclic compound epoxyethane

by air
$$\left(\begin{array}{c} CH_2 = CH_2 \xrightarrow{Ag} CH_2 \\ \hline [O] CH_2 \end{array}\right)$$
, polymerize with various catalysts or under pressure, used as final.

catalysts or under pressure, used as fuels.

- Alkoxide. An organic compound containing an ion of the type RO; where R is an alkyl group, obtained by the reaction of metallic sodium on alcohol. Example: sodium ethoxide
- Alkyl. Residue left when a hydrogen alom is removed from an aliphatic hydrocarbon, e.g., CH₃, C₂H₅, etc., known as free radicals and used as intermediates in many reactions such as the thermal degradation of alkanes and petrochemical process.
- Alkylation. A process involving the introduction of alkyl groups into hydrocarbon chains or aromatic rings, can be effected thermally, the production of branched-chain hydrocarbons by reacting alkeres with iso-alkanes, e.g., the production of iso octane as a high anti-knocking gasoline by reacting iso-butane
- Alkynes, Acetylenes. Compounds containing C groups. Thus methylacetylene (CH₃C\equiv CH) is propyne; dimethylacetylene,

Allantoin, Glyoxylidiureide, C₄H₆N₄O₃. M.P. 235°—236°C. An end product of purine metabolism, present in the urine of mammals.

- Allenes, 1, 2-Dienes. Unsaturated organic compounds with general formula $C_nH_{2^{n}-2}$ and skeleton >C=C=C<, derivatives of 1, 2-propanediene, undergo typical reactions of alkenes with, e.g., hydrogen, halogens and hydrogen halides, colourless liquids with garlic odours but higher members solids. prepared from 1, 2, 3-tribromopropane in the presence of KOH and Z_n/C_2H_5OH .
- Allethrin, C₁₉H₂₆O₃. A viscous liquid, shows toxicity to many insects such as house fly, used mainly as the synthetic material.
- Allobarbitone, Diallyl Barbituric Acid, C₁₀H₁₂N₂O₃. White powder having hypnotic properties like barbitone, M.P. 172°—174°C.
- Allomone. A defensive secretion, a floral scent attracting pollinating insects.
- Allorhanic Acid, Carbamyl Carbamic Acid, C₂H₄N₂O₃, NH₂CONH COOH. Not available in free state but as NH₄, Ba, Ca, K and Na salts, prepared by passing cyanic acid into alcohols or a solution of alcohol or phenol in benzene.
- Allostearic Effects. The control of working of some enzymes by their interaction with small molecules at sites distant from the active sites.
- Allotropy. The existence of an element in more than one physical state, show difference in crystals (structures), e.g., graphite and diamond or white tin and grey tin, etc., various forms termed as allotropes.
- Alloxantin, C₈H₆O₄N₂. Prismatic crystals with two molecules of water, sparingly soluble in water, decomposing at 253°-255°C, prepared by oxidising uric acid or reducing meso-oxalylurea

of hydrolysis of glycosides convicium from soyabeans.

- Alloy. A compound or association of two or more metals and one or more non-metallic elements, (e.g., bronze or brass and steel), completely homogeneous mixture or may contain small particles of one phase in the other phase, Alloying is one of the common methods of increasing useful properties of metals, e.g., hardness, strength, resistance to corrosion.
- Allyl Rearrangement. A process involving the migration of a double bond in a three carbon system from C atoms one and two to C atom two and three and the substituent simultaneously migrates from C atom 3 to C atom 1.

$$C^1 = C^2 - C^3 - X \rightleftharpoons X - C^1 - C^2 = C^3$$

Example: The hydrolysis of crotyl chloride (2-butanyl-chloride to form crotyl alcohol,

$$CH_{3}CH=CH-CH_{2}CI$$

$$CH_{3}. CH. CH=CH_{2}$$

$$CH_{3}. CH CH=CH_{2}$$

$$CH_{3}. CH=CH. CH_{3}.OH.$$

$$CH_{3}. CH=CH$$

- Allyl Polymers. Polymers derived from monomers containing the CH₂=CH-CH₂—group (allyl, propylene groups), e.g, Allylphosphates make polymers with good flame resistant properties, allylethers of carbohydrates and other polyols together with styrene and other vinyl polymers used as adhesives, and furniture finishes.
- Alpha Particles, α·ray. A heliun ion (He²⁺) or a doubly ionized He atom, emitted with high velocity (C.2×10⁹Cm.Sec⁻¹) by without acceleration as bombarding agents in nuclear disintespecificity of the source of emission.
- Alum, Potash Alum. KAl (SO₄)₂. 12H₂O. A substance crystallizes in large colourless octahedra, crystals convert white on exposure to air, dehydrated at a duel red heat to form burnt mixing equimolecular quantities of solutions of ammonium a mordant for dyes, water proofing agent and as a tanning
- Alumina, Al₂O₃. White powder insoluble in water, amphoteric nature, occurs naturally as bauxite, corundum and white sapphire, manufactured by heating aluminium hydroxide, used (corundum), in furnace lining and as a catalyst.

Alumina Gel. See aluminium hydroxide.

Aluminates. See aluminium hydroxide.

- Aluminium, Al. At. No. 13, At. wt. 26.9816, M.P. 660.37°C, B.P. 246.7°C, D 2.702, occurs as silicates and as the hydrated oxide, manufactured by electrolysis of Al₂O₃ dissolved in Ma₃AlF₆ (cryolite) with added Li₂CO₃ using carbon electrodes, pure Al obtained by electrolysis of NaAlF (C₂H₅)₃, pure Al unattacked by water, corroded by salts of sea water, reduces many oxides (Goldschmidt, thermite reaction), dissolved by acids; Al and its alloys used in aircrafts, utensils, overhead cables, as a catalyst, the oxides used in, refractories, abrasive, cements, ceramic and gemstone, etc. Al salts also used in water purification.
- Aluminium Bromide, AlBr₃. A white solid soluble in water and many organic solvents, formed from Al and Br₂, forms hydrates.
- Aluminium Chemistry (A1). An element of III group, electronic configuration 1s²2s²2p⁶3s²3p³, E°-1 66 volt (Al³⁺-→Al in acid solution).
- Aluminium Chloride, AlCl₃. Colourless or yellowish hexagonal crystals, D 2.44, B.P. 180°, M.P. 192.6° at 1715 mm., soluble in water, prepared by passing hydrogen chloride or Cl₂ over heated aluminium, or chlorine over heated aluminia and carbon. Forms double salts (e.g., 2AlCl₃, 2, 3 or 6 NaCl), addition compounds (e.g., AlCl₃, H₂S; AlCl₃NOCl) and aluminates. Solid AlCl₃ contains octahedrally coordinated Al, dimeric [Cl₂Al (μ-Cl)₂ AlCl₂] in the gas phase. AlCl₃ is used in the cracking of oils and as a catalyst (e.g., Friedel-Crafts reaction).

Aluminium Ethoxide, Aluminium Ethylate, (CH₃CH₂O)₃ Al. M.P. 139°C, B.P. 320°C.

Aluminium Fluoride, AIF₃. A white crystalline solid, soluble in water but insoluble in most organic solvents, used as an additive to the cryolite electrolyte in the production of Al, D 2.88—3.22.

Aluminium Hydroxide. Al (OH)₃. A white powder prepared as a colourless gelatinous precipitate by adding ammonia solution to a solution of Al salts. The gelatinous "gels" of Al (OH)₃ partially dried at 200-500°C, are valuable drying agents, catalysts and absorbents (alumina gel). Aluminium hydroxide is amphoteric, dissolving in acids and in alkalis.

Aluminium Iodide, All3. Formed from Al and I2.

Aluminium Nitrate, $Al(NO_3)_3$. nH_2O , where n=9.

Aluminium Organic Derivatives, Aluminium Alkyls. Formed Al plus R₂Hg, AlCl₃ plus RMgX, or on a commercial scale, Al-H plus C_nH₂ⁿ (alkene) to produce AlC_nH₂ⁿ+1. All Al organic

derivatives are easily oxidised. The lower alkyls are polymeric liquids and spontaneously inflammable. Complexes AlR₃L, are readily formed, used for introduction of alkyl groups, polymerization. (Ziegler process) and telomerization of alkenes to medium chain derivatives for detergents and fats.

Aluminium Oxide. (See Alumina).

- Aluminium Oxy-acid Salts. Soluble in water (except the phosphate) and, generally, form hydrolysed solutions, heavy hydrated salts. Important salts are the ethanoate (acetate), the nitrate (used as a mordant) and the sulphate (used in water treatment and paper sizing).
- Aluminium Silicates. Al₂SiO₅ (cyanite, sillimanite and andalusite) contains discrete SiO₄⁴⁻ tetrahedra.
- Aluminium Sulphate, Al_2 (SO₄)₂. nH_2O (n=0, 6, 10, 16, 18, 27), occurs in nature as alumite.
- Aluminoferric. Impure Al₂(SO₃)₃ having some Fe¹¹ and Fe¹¹¹, used in water and sewage purification.
- Aluminon, $C_{22}H_{23}N_3O_9$. An organic reagent used for the detection and estimation of Al. A brownish red powder, soluble in water, forms a bright red lake with Al which can be estimated colorimetrically, also used for the detection of scandium and indium.
- Aluminosilicates. Silicates containing Al, have Al in tetrahedral position replacing silica, an M⁺ cation must be present for balancing the charge for every Si replaced by Al, Al also present in octahedral co-ordination in silicates when it functions as a normal cation. Examples: Feldspars, mica, zeolites and some clay materials.
- Aluminium Cement. Made up of calcium aluminate, resembles
- Alums. A group of crystalline double sulphates with general formula M^IM^{III} (SO₄)₂, 12H₂O. The sulphate radical may be replaced by SeO₄²⁻, BeF₄²⁻ or ZnCl₄²⁻, KAl (SO₄)₂. 12H₂O is known as potash alum.
- Alundum. An artifical variety of corundum (a·Al₂O₃), formed by lusing calcined bauxite in an electric furnace, and allowing the molten product to cool rapidly. Used for highly refractory bricks, crucibles and muffles, etc.
- Alunite (or alumstone). A mineral composed of a hydrated basic sulphate of K and Al, [KAl₃ (SO₄)₂ (OH)₆]; the potassium may be partially replaced by sodium. Used commercially as a source of potash alum and of potassium sulphate, sp. gr. 2.75.
- Am, Americium. A highly toxic radioactive element, synthesised from plutonium, M.P. 994°C, B.P. 2607°C, R.I. 13°67, stablest lsotope²⁴³ Am $(t_{1/2} 7.4 \times 10^3 \text{ years})$.

Amino Acids 25

Amalgam. An alloy of mercury with one or more than one metals, may be a liquid or solid, Na/Hg amalgam is used as a source of nascent hydrogen.

Americium. See Am.

Aminitius. A group of very toxic cyclic peptides present in the mushroom Amanita phalloides.

Amaranth. An important red 930 dye used in food stuffs.

Amatol. A mixture of NH4NO3 and TNT used as explosive.

Ambident Ligands. Ligands which can use more than one coordination site, e.g., NH₂. CH₂COOH (co-ordination through N or O).

Amblygonite, Li Al (F, OH) PO4. A lithium ore.

Ametryne, C₉H₁₇N₅S. A soil-acting herbicide, a triazine derivative, forms colourless crystals, M.P. 84-86°, used as a pre-and postemergence selective herbicide to control, broad-leaved and grass weeds in crops.

Amicron. Particles too small to be visible even in the ultramicroscope (Zsigmondy).

Amides. Organic compounds with general formula RCONH₂, white solids formed by the partial dehydration of the ammonium salts of a carboxylic acid:

RCOO-NH4+-H2O-RCONH2

Amides form carboxylic acids and nitriles (Hofmann degradation). Inorganic amides contain the ion NH₂⁻ and formed by the action of ammonia on metals or by the amonolysis of nitrides, heavy metal amides are, generally, explosive.

Amines. Organic compounds derived from ammonia by replacement of one or more of its hydrogen atoms by hydrocarbon groups. Replacement of one, two and three hydrogen atoms forms primary, secondary and tertiary amines respectively. Primary amines are obtained by reduction of nitro-compounds, nitriles, ketoximes and amides, or by treating amides with NaOBr. Secondary amines are obtained by reducing Schiff's bases. A general reaction for the preparation of amines is that between petassium phthalimide and a halogen compound, with subsequent hydrolysis of the product.

Aminoacetal, C₆H₁₅NO₂, H₂N.CH₂.CH(OC₂H₅)₂. Colourless oily liquid with smell resembing ammonia, B.P. 172°-174°C.

Amino Acids. Organic compounds containing both the carboxyl, COOH and the amino, NH₂, group, e.g. glycine (H₂N . CH₂. COOH). Obtained from proteins and are colourless crystalline substances which melt with decomposition.

- Aminoazobenzene, $C_{12}H_{11}N_3$, N=N-N-N+1.
 - Brownish yellow needles, M.P. 127°C. The hydrochloride produces steel blue needles. Used as a first component in the preparation of azo-dyes.
- 4-Amino Benzoic Acid (p-Aminobenzoic Acid), P.A.B. C₇H₇O₂N. Yellowish red crystals, soluble in water, M.P. 187°C and a member of vitamin B group.
- Aminocaproic Acid, $H_2N \cdot N(CH_2)_5 \cdot COOH$, $C_6H_{13}NO_2$. Antifibrinolytic agent used to treat thrombosis in the deep veins.
- 5-Aminolaevulinic Acid, C₅H₉NO₃, H₂NCH₂COCH₂CH₂COOH. A basic unit for the biosynthesis of porphyrins (e.g., provide all the C and nitrogen atoms in the haem of haemoglobin, myoglobin, etc.). Also present in the corrin ring of vitamin B₁₂.
- Aminophyllius. Theophylline (75-82%) with ethylenediamine (12-14%). Used as a diuretic and as a bronchodilator in the treatment of asthma.
- Aminoplastics and Resins. A group of plastic materials obtained from resins derived from amino compounds such as urea, melamine, thiourea and aniline by condensation with form aldehyde.
- 4-Aminosalicylic Acid, P.A.S., C₇H₇NO₄. An important drug used to treat tuberculosis infections.
- Amines. Complexes containing ammonia molecules coordinated to a metal ion; e.g., [Cu(NH₂)₄]²⁺.
- Ammonal. An explosive of a mixture of ammonia, nitrate, nitrotoluene, Al powder and charcoal.
- Ammonia, NH₃. A colourless gas with a characteristic pungent smell. Readily liquefied by cooling or compression, B.P.—33.5°C and M.P.—77.7°C. Soluble in water, a solvent for organic compounds but a poor solvent for inorganic compounds. Detected by Nessler's reagent and blackening of Hg₂(NO₃)₂.
- Ammonical Liquor. An aqueous liquid containing admixed tarry matter and organic compounds.
- Ammonium Bicarbonate. See ammonium hydrogen carbonate.
- Ammonium Bromide, NH₄Br. A colourless crystalline solid with saline taste. Soluble in water, sublimes on heating and turns yellow in air.
- Ammonium Carbonate, $(NH_4)_2CO_3$. ("Sal Volatile"). A double salt of ammonium hydrogen carbonate and ammonium carbonate having formula NH_4HCO_3 , NH_2COONH_4 . Manufactured by heating a mixture of chalk and ammonium chloride or sulphate.

- Soluble in water and decomposes to NH₃, CO₂ and H₂O on heating. Used in baking powder, smelling salts, dyeing in wool treatment, etc.
- Ammonium Chloride, NH₄Cl. ("Sal Ammoniac"). A white crystalline solid with saline taste. Soluble in water, sublimes on heating, used in dry cells, as a mordant, and as a soldering and galvanizing flux.
- Ammonium Chromate, $(NH_4)_2CrO_4$. A golden yellow solid, soluble in water and decomposes on heating to the dichromate, NH_8 . and H_2O .
- Ammonium Dichromate, $(NH_4)_2Cr_2O_7$. A red crystalline solid, very soluble in water and decomposes on heating to N_2 , H_2O , and Cr_2O_5 .
- Ammonium Hexachlorostannate (Pink Salt), (NH₄)₂SnCl₆. An important mordant.
- Ammonium Hydrogen Carbonate, Ammonium Bicarbonate. NH4HCO3. A white crystalliue solid. A component of commercial ammonium carbonate and produced by decay of nitrogenous materials and present in guano. Used in baking powder and medicines.
- Ammonium Hydroxide, (NH₄OH). An aqueous solution of NH₃, a weak base and forms salts with acids. Forms two hydrates on freezing of aqueous solution of ammonia (e.g., NH₃ H₂O and 2NH₃. H₂O).
- Ammonium Iron (III) Sulphate, (NH₄)₂ FeSO₄. 6H₂O (Mohr's Salt). A double sulphate of iron and ammonium, light green crystalline solid soluble in water.
- Ammonium Molybdate. A range of salts obtained from NH₃ and molybdic acid. The commercial salt has the formula (NH₄)₆Mo₇O₂₄,4H₂O.
- Ammonium Nitrate, NH₄NO₃. A colourless crystalline solid, soluble in water, sublimes (NH₃ and HNO₃) and on gentle heating decomposes to N₂O, used as an explosive and as a fertilizer.
- Ammonium Perchlorate, NH₄ClO₄. A white crystalline solid, used in propellants.
- Ammonium Phosphomolybdate, $(NH_4)_3Mo_{12}PO_{40}$, xH_2O . A bright yellow ppt obtained from a phosphate, ammonium molybdate and HNO₃. Used as a test of phosphate.
- Ammonium Sulphamate. A translocated and soil acting herbicide.
- Ammonium Sulphate, $(NH_4)_2SO_4$. A colourless crystalline solid, soluble in water. Forms N_2 , NH_3 , SO_2 and H_2O , on strong heating. Used as a fertilizer.

- Ammonium Sulphides, (NH₄)₂S (NH₃ and H₂S at -18°C) and NH₄HS (NH₃ and H₂S in CH₃OOC. CH₃ solution) are relatively unstable. A solution of the hydrosulphide prepared by passing H₂S through strong aqueous ammonia is used in qualitative analysis as "yellow ammonium Sulphide"
 - Amorphous. A solid substance which is not crystalline (no ordered arrangement of atoms, ions and or molecules). Glasses and super-cooled liquids are amorphous while many amorphous substances have microcrystalline structure, e.g., charcoal, coke and soot.
 - Ampere, A. The SI base unit of electric current.
 - Amperometric titrations. A method of analysis in which current passing through an electrolytic cell is plotted against added titrant. End points are represented by sharp breaks in the curves.
 - Ampholyte. A substance which can behave both an acid and a base.
 - Amphoteric Oxide or Hydroxide. An oxide which can function both an acid and a base, e.g., ZnO reacts with acids to form zinc salts and in alkalis to form zincates [Na₂Zn(OH)₄]. Al(OH)₃ is an amphoteric hydroxide.
 - amu, atomic mass unit. Units of atomic weight in terms of an arbitrary standard (12C).
 - Amylacetate, C₇H₁₄O₂. A colourless volatile liquid with a strong pear-like smell, soluble in alcohol and ether. Used as a solvent for cellulose acetate lacquers and paints; an alcoholic solution is used as "essence of Jargonella pears".
 - Amyl Alcohol, C₅H₁₂O. An aliphatic alcohol with eight isomeric alcohols. Complex mixtures of amyl alcohols are manufactured from chloropentanes derived from petroleum.
 - Amylases, diastase. Enzymes which break down starches and glycogen to the maltose stage. Occur in animal and plant tissues.
 - Amylopectin. See starch.
 - Anabolic Agents. Drugs which promote storage of protein and stimulate tissue metabolism, (e.g., testosterone).
 - Anaesthetics. Compound used to remove the perception of pain and other stimuli. Local anaesthetics act at only the site of application by freezing, e.g., ethyl chloride, or by affecting the nerves, e.g., Cocaine, while general anaesthetics produce total anaesthesia, e.g., halothane, N₂O.

Analgesics. Compounds which can reduce pain perception, e.g., morphine.

Analysis. The process for determining the constituents or components of a sample. Two major classes of analysis are: Qualitative analysis, in which the identity of an unknown substance is to be determined and quantitative analysis, in which the amount of a substance present in a mixture of substances or solution is determined (e.g., volumetric and gravimetric analysis).

Anatase, T1 O2, A steel blue or yellow form of TiO2.

Anation. Replacement of an uncharged ligand e.g., H₂O, in a complex by Cl⁻ ion.

Andrews Titration. A titration for estimation of reducing agents.

Angstrom Unit, (A°). A unit of length defined as 10^{-8} cm, (10^{-10}m) . Used for expressing wave length of light and molecular and lattice dimensions.

Anhydride. A compound formed by elimination of one or more molecules of water from one or more molecules of an acid or, a base e.g., CO₂ is the anhydride of H₂CO₃ and phthalic anhydride

Anhydrite. Mineral CaSO₄ generally associated with some gypsum, CaSO₄, 2H₂O, and rock salt. Used in the commercial preparation of H₂SO₄ and ammonium salts.

Anhydrone. A name for magnesium perchlorate.

Anilde. Acyl derivatives of aniline.

Aniline, C₆H₅NH₂ (aminobenzene). A colourless oily liquid formed by reducing nitrobenzene, used in the manufacture of antioxidants, for dye preparation and in the rubber industry.

Anion. A negative charged ion, formed by addition of electrons to atoms or molecules, e.g., NO₃, Cl⁻ or SO₄²⁻ ions.

Anisotropic. A substance (crystal) showing different physical properties (e.g. refractive index, thermal conductivity) in different principal directions. All crystalline systems, except cubic system are anisotropic.

Annealing. The process of heating and cooling of a substance toproduce a ordered structure of the substance.

- Anode. The electrode of a electrochemical cell and carries a positive charge. In a discharge tube or electronic device, the anode is a terminal at which electrons flow out of the system or negative ions are discharged.
- Anodic Oxidation. The process of oxidation taking place at anode in an electrolytic cell.
- Anodizing. Process for protecting metals from corrosion by anodic oxidation. Commonly used for protecting Al with an oxide layer formed in an electrolytic cell. The layer of Al₂O₃ is porous and can be coloured with certain dyes.
- Anomers. A specific term used to explain carbohydrate stereoisomers e.g., α -D-Glucose and β -D-Glucose are α -and β -anomers of glucose.

Antabuse. A drug used in the treatment of alcoholism.

Antacids. Substances used to reduce the amount of acid in the stomach.

fluorescence, present in coaltar.

Anthracite. A bright hard coal containing more than 93% carbon,

Antibiotic. An organic substance produced by micro-organisms capable of inhibiting the growth of, or destroying another micro-organism. Isolated chiefly from bacteria (e.g., (from Fungi).

Antibodies. Substances present in the serum are known antibodies. Protein molecules present in the serum which are produced in the body in response to the presence of forcign substances of the immunology.

Antibonding Orbitals. See molecular orbitals.

Anticathode. See X-ray tube.

- Antifoaming Agents. Substances which prevent the formation of foams, e.g., polyamide and octanal.
- Antifreezing Agent. A material used to prevent freezing in the cooking system of internal combustion engines, e.g., ethylene
- Antigens. Macromolecular proteins which on introduction in to the blood of animals stimulate the production of antibodies, e.g., polysaccharides and nucleic acids.

- Anti-knock Additives. Substances which inhibit the knocking property of gasoline, e.g., a mixture of lead alkyls with di-bromoethane and dichloroethane (ethyl fluid) acts as an antiknock when added to gasoline.
- Anti-knock Value. See octane number.
- Antimony, Sb. At. No. 51, At. wt. 121.75, a toxic element existing in two allotropic forms. Occurs as stibnite (Sb₂S₃) and is used in alloys and as a semiconductor.
- Antimony Chloride (antimony trichloride, SbCl₃). A white deliquescent solid. Readily hydrolysed by cold water to form a white ppt. of antimony (III) chloride oxide (antimony chloride): SbCl₃+H₂O -SbOCl+2H₂O.
- Antimony (III) Oxide, Sb₂O₃. A white amphoteric oxide with a strong tendency to act as a base.
- Antimony (V) Oxide, Sb₂O₅. A yellow solid and an acidic oxide, slightly soluble in water.
- Antimony Potassium Tartrate, Tartar Emetic, KsbO (C4H4O6), 1H2O. A white crystalline solid used as an emetic.
- Antioxidants. Substances which slow the oxidation rate in autooxidizable substances, e.g., substituted phenols, aromatic amines and sulphur compounds. Used for the protection of fats, preservatives, inhibitors in petroleum industry.
- Antiparticle. An elementry particle with charge properties opposite to those normally found, e.g. positive electron.
- Antipyretics. Drugs used to reduce temperature in the case of fever e.g., phenazone.
- Antiseptics, Substances (or materials) which can stop the growth of micro-organism, if applied to living tissues, e.g., iodine.
- Antitoxins. Antibodies formed in blood to which bacterial toxins have been introduced. A given antitoxin will neutralize its own toxin, with which it apparently combines in a quantitative
- Apatite, Ca5F(PO4)3. A "mineral" phosphate used for the manufacture of phosphorus compounds, e.g., Superphosphates con-
- Aprotic Solvents. Inert solvents.
- Aqua Regia. A mixture of concentrated HNO3 and three to four parts of HCl. Dissolves all metals such as gold and platinum due to the formation of nitrosyl chloride (NOCI) and chlorine.
- Ar. Argon. An inert gas.
- Aragonite, CaCO₃. A variety of calcium carbonate.
- Arc Spectrum. Emission spectrum formed by the excitement of a

π-Arene Complexes. Complexes in which aromatic system is bonded to a metal through π-electron.

Argentite, Ag₂S. An important ore of Ag.

Arginase. An enzyme.

Arnd's Alloy. An alloy of Cu (60%) and Mg (40%) used for the estimation of nitrates.

Arndt-Eistert Synthesis. A process of converting a carboxylic acid to its next higher homologue, or to a derivative of a homologous acid, e.g., amide.

Aromatic. A characteristic (promaticity) that differentiates the benzene derivatives from cuphatic and alicyclic compounds.

Aromatic compounds contain benzene nucleus and un-

dergo electrophilic substitution reactions.

Arrhenius Equation. An equation co-relating the rate constant of a chemical reaction and temperature at which reaction takes place.

$$k=A \exp \left(-\frac{E_{\sigma}}{RT}\right)$$

where k is the velocity constant, A and E_a are constants, known as pre-exponential factor and the activation energy of the reaction. T is the thermodynamic temperature and R is the gas constant. Used in determining the activation energy of a reaction, in processes such as diffusion, electrolytic conduction.

Arsenic, As. M.P. 817°C, a toxic metalloid element existing in several allotropic forms; brittle grey is the most stable form. Belongs to V group of periodic table and used in medicines and insecticides (as poison).

Arsenic Chloride, AsCl₃. A poisonous oily liquid, fumes in moist air. Covalent with non-metallic properties.

Arsenic (III) Oxide, As₂O₃. A colourless, crystalline and poisonous solid. Occurs as a dimerized structure (As₄O₆), and a basic oxide.

Arsenic (V) Oxide, As₂O₅. A white amorphous deliquescent solid, an acidic oxide.

Arsine, AsH₃ (Arsenic Hydride). A poisonous colourless gas with bad smell. Produced in the analysis of arsenic (Marsch's test).

Aryl Group. An organic group obtained by removing hydrogenatom from an aromatic hydrocarbon.

- Arsenides. Compounds obtained from metals and arsenic.
- Arsine, AsH₃, (Arsenic Hydride). A poisonous colourless gas with bad smell. Produced in the analysis of arsenic (Marsch's test).
- Asbestine. A fibrous species of talc mixed with tremolite asbestos, used in paints and as rubber fillers.
- Ascorbic Acid, Vitamin C, C₆H₈O₆. Extracted from plant source such as rose hips and citrus fruits. Essential constituent in the formation of collagen, bone and teeth and for healing wounds. Used as a developer in photography (alkaine).
- Aspargine, Aminosuccinamic Acid, C₄H₈N₂O₃. An amino acid widely present in plants, in all Leguminosae and Gramineae and in many seeds, roots and buds.
- Aspartic Acid, Aminosuccinic Acid, C₄H₇NO₄. One of the acidic amino acids obtained by the hydrolysis of proteins.
- Asphalt. A natural mixture of bitumens, e.g., rock asphalt.
- Aspirin, o-A-Acetylsalicylic Acid, C9H8O4. An analgesic and antipyretic substance.
- Associated Liquids. Liquids having combined aggregates of two or more molecules, e.g., water $(H_2O)^n$ and alcohol show high boiling points.
- Astatine At. M.P. 302°C, B.P. 337°C. A radioactive element belonging to halogen group. Occurs in small quantities in uranium, 20 small time span isotopes are known. Stablest Isotope 210 At. $t_{1/2}$ 8'3 hours).
- Asymmetry. A property of lacking symmetry, in molecular or of crystal structure, e.g., central C atom in tartaric acid is asymmetric.

Asymmetry leads to optical active nature of compounds. Even some crystals show asymmetry, e.g., quartz.

- Atmolysis. The separation of gases by using their different rates of diffusion (Graham's law of diffusion of gases).
- Atmosphere. A unit of pressure defined as 101325 pascles (atmospheric pressure).
- Atom. The smallest part of an element which can exist as a stable entity. Consists of a nucleus (protons-f-neutrons) and surrounding electrons.

- Atomic Absorption Spectroscopy. An analytical technique involving the absorption of characteristic radiation by non-excited atoms in vapour phase.
- Atomic Emission Spectroscopy. An analytical method employed for the determination of traces of metals. The sample is vaporized and excited in an arc or flame and the emission spectra recorded.
- Atomic Energy. According to Einstein's relativity theory, the mass and energy are interconvertible, and are related by the equation. $E=mC^2$, where E (ergs) is the energy associated with mass m (gms), and c is the velocity of light $(3\times10^{10}$ cm. per sec). Energy released in transmutation process is known as atomic energy.
- Atomic Heat. Atomic heat of an element is the product of its atomic weight and specific heat. It is the heat capacity, water being taken as unity, of one gram-atom of the element.
- Atomicity. The number of atoms per molecule of a compound, e.g., CH₄ has an atomicity of five.
- Atomic Mass. See Atomic weight.
- Atomic Number. The number of unit positive charges (protons) present in the nucleus of an atom of the element. Determines the position of the element in the periodic table. Determined by X-ray spectrum of the element.
- Atomic Orbital. The region around an atomic nucleus in which there is a high probability of finding an electron. Four atomic orbitals are: s, p, d and f containing 2, 6, 10 and 14 electrons in each orbital respectively.
- Atomic Radii. Half the closest distance of approach of atoms in the structure of elements. The atomic radius of an element depends upon the nature of the interatomic forces holding the atoms together and also the number of neighbours around each atom, e.g., in the two forms of carbon the C—C distances are 1.54 A°(in diamond) and 1.42 A° (in graphite).
- Atomic Spectrum. See line spectra.
- Atomic Weights. Defined relative to ¹²C as 12.00, and are ratios of masses. Also called formerly as atomic weights.
- Atrazine, C₈H₁₄ClN₅. A soil acting herbicide, colourless, M.P. 173-175°C.
- Atropine, (±)-hyoscyamine, C₁₇H₂₃NO₃. M.P. 116°C, used to dilate the pupil.

Au. Gold.

Aufbau Principle. A principle that controls the order in which the electrons in an atom or molecule are filled up in the orbitals. In other words, explains the building up of the electronic configuration of an atom or molecule in its ground state, the electrons are filled in the orbitals in order of increasing energy. The order is as follows:

 $1s^2$, $2s^2$, $2p^6$, $3s^2$, $3p^6$, $4s^2$, $3d^{10}$, $4p^6$, $5s^2$, $4d^{10}$, $5p^6$, $6s^2$, $4f^{14}$, $5d^{10}$, $6p^6$, $7s^2$, $5f^{14}$, $6d^{10}$.

Auric. Gold (III) Compounds, e.g., AuCl3.

Aurous. Gold (I) Compounds, e.g., AuCl.

Austemite. See Iron, steel.

Auto-catalysis. A process in which one of the products catalyses the further reaction, e.g., Mn²⁺ ions autocatalyse the KMnO₄ and oxalic acid interaction.

Autoclave. Metal equipment used for reaction between gases under pressure and liquid and solid reactants.

Autolysis. A process of self-destruction which occurs in living systems after death by enzymes released from lysosomes.

Autoxidation. A process of oxidation which occurs only when another oxidation reaction is occurring simultaneously in the same system, e.g., the oxidation of oil of turpentine by atmospheric oxygen. When this reaction is taking place it is possible to oxidise KI to I_2 .

Autunite, Ca(UO2)2(PO4)2, nH2O. A mineral of uranium.

Auxins. Plant hormones.

Auxochrome. Used to modify the colour of a chromophore.

Aviation Turbo-fuels. Fuels used for air craft jet engines.

Avogadro's Number, N_A. The number of particles in one mole of a substance. Its value is 6.02252×10²³ mol⁻¹.

Avogadro's Law. Equal volumes of all gases under similar conditions of temperature and pressure contain equal number of molecules (strictly applicable to ideal gases).

Axial Ratio. The ratio of the cell dimensions (a, b and c) in a crystal; b being taken as unity. In a cubic system a=b=c. In hexagonal and tetragonal systems a=b≠c. In triclinic, monocinic and orthorhomic systems a≠b≠c and axial ratios

are
$$\frac{a}{b}:1:\frac{c}{b}$$
.

Axis of Symmetry. See crystal symmetry.

- Azeotrope (Azeotropic Mixture). Mixture of liquids which distile with constant composition, i.e., the composition of liquid phase is same as that of the vapour phase. Azeotropes are not chemical compounds as their composition and boiling points vary with pressure. They can be broken by distillation in the presence of a third liquid, by chemical reactions, adsorption and fractional distillation. Many liquid azeotropes show a minimum boiling point, (e.g., methanol and chloroform) whereas others show a maximum B.P. (e.g., hydrochloric acid and water) when vapour pressures are plotted as a function of change in the composition of the mixture.
- Azeotropic Distillation. A process used to separate mixture of liquids that can not be separated by simple distillation (azeotropic mixtures). The two substances may also have similar boiling points. The process of azeotropic distillation requires the addition of a third solvent to form a new azeotrope with one of the components and this is then removed and subsequently separated in a separate column, e.g., the dehydration of 96% ethanol to absolute alcohol (100%) using benzene.
- Azide. An inorganic compound containing the ion N₃⁻ and an organic compound of general formula RN₃. Salts of hydrazoic acid, HN₃, e.g., NaN₃. The azide group is a good complexing agent.
- Azimuthal Quantum Number, I (angular momentum quantum number). Have values 0 to (n-1), where n is the principal quantum number.
- Azine. An organic heterocyclic compound possessing a hexagonal ring containing nitrogen and carbon atoms, e.g., Pyridine. Also used for compounds containing multiple N₂ groups.
- Azobenzene, C₁₂H₁₀N₂, Ph.N=N.Ph. Orange red crystals, M.P. 68°C.
- Azo Compounds. A large group of compounds containing the -N=N-chromophere attached to two aromatic nuclei and which absorb light in the visible range of the spectrum, e.g.,

benzeneazophenol (N=N- N-OH).

- Azo Dyes. Organic compounds containing one or more—N=N—groups. Used as pigments, as dye-stuffs in colour photography.
- Azine Dyes. Water-insoluble dyes also known as ice colours or ingrain dyes.
- Azurite, Cu₃ (OH)₂(CO₃)₂. Blue basic carbonate of copper, used as an artist's colour.

B. Boron.

Ba. Barium.

Bacitracin. Antibiotic produced by a strain of B. subtilis. Used against organisms resistant to penicillin, chiefly for skin and eye infections.

Back Bonding. Overlap of a filled orbital (d orbital), of an acceptor with a vacant orbital, d, p or π (bonding or anti-bonding) of a ligand, e.g., used in metal carbonyls formation.

Back e.m.f. An e.m.f. that opposes the normal flow of electric charge in a circuit.

Baddeleyite. Principal mineral ore of zirconium (Zr).

Bakelite. An important class of phenol-formaldehyde/plastics and resigns.

Baking Powders. Substances used in baking and substitutes for yeast, e.g., potassium hydrogen tartrate or sodium hydrogen phosphate with NaHCO₃ or NH₄HCO₃ together with flour or starch (inert material).

BAL (British Anti-Lewisite), C₃H₈OS₂. An oil, M.P. 77°C, used as an antidote to poisoning by organic arsenicals, Hg, Cu, Zn and Cd but not Pb.

Balance. An instrument used for weighing.

Ball Mill. A device commonly used in the chemical industry for reducing the size of the solid material.

Balmer Series. A series of lines present in the spectrum of radiation emitted by excited hydrogen atoms. The lines correspond to the atomic electrons falling into the second lowest energy level, emitted energy as radiation. The wavelength (λ) of the radiation in Balmer series are expressed by the equation

$$\frac{1}{\lambda} = R\left(\frac{1}{z^2} - \frac{1}{n^2}\right)$$

where n is an integer and R is the Rydberg constant.

Band Spectrum. A spectrum produced as a number of bands of emitted or absorbed radiation. Band spectrum is a characteristic of a molecule. It consists of well-defined groups or bands, of closely packed lines. The close lines seen under high power microscope are the results of different vibrational states of molecules.

- Band Theory of Solids. A theory involving the quantum mechanical treatment of electrons (delocalized electrons) which give rise to the band theory of solids.
- Barbitone, C8H12N2O3. A barbituric acid derivative used as a sedative.
- Barbituric Acid, Malonyl Urea, C4H4N2O3. M.P. 253°C. A dibasic
- Barft Process. A process used for protecting iron by heating it in steam to form a layer of Fe₃O₄.
- Barium, Ba. A dense metal, M.P. 714°C, B.P. 1537°C, a member of 11 group of the periodic table and a typical alkaline earth metal. Chief ores are: witherite (BaCO₃) and barytes (BaSO₄), Its salts are highly toxic. Used in paints (as BaSO₄, Lithopone), rat poisoning (as BaCO₃), X-ray diagnosis (as Ba meals) and in
- Barium Carbonate, BaCO₃. A white crystalline solid. Occurs as witherite and used as a rat poison.
- Barium Diphenylamine-p-Sulphonate. A redox indicator used in
- Barium Halides, BaX2. Four halides are known, e.c., BaF2, BaCl2 (forms BaCl₂. 2H₂O; used in the electrolytic production of Ba),
- Barium Hydroxide, Ba(OH)2, Baryta. White hydroxide used in volumerric analysis, as a plastic stabilizer, in pigments, gasoline additive and as a vulcanization acceptor.
- Barium Oxides. Two oxides are known, e.g., BaO and BaO2 (barium peroxide). BaO₂ is used as a bleaching agent in hydrogen
- Barium Sulphide, BaS, Black Ash. Strongly phosphorescent.
- Barrel. A unit of measuring volume in petroleum industry (one
- Base. A substance which produces free hydroxyl ions (OH- ions) in aqueous solution, e.g., NaOH, Ba(OH)₂ and Na₂CO₃ solutions. pH more than 7.00.
- Base Metal. A metal such as ion, copper, distinguished from a
- Basic Salt. A compound intermediate between a normal salt and a hydroxide or oxide. Formed by hydrolysis of metal salts and generally precipitated by sodium carbonate solution, e.g.,
- Basic Slag. See slag.

Bastenaesite. A mineral, CeFCO3, used for the preparation of lighter lanthanides.

Batteries. A number of electric cells, working together which produce electricity, e.g., the Leclanche's Cell (MnO, \rightarrow Mn³⁺ \rightarrow Mn²⁺. $||Zn \rightarrow Zn^{2+}|$ in a NH₄Cl-MnO₂-Zn system).

Bauxite. A principal ore of aluminium, used in the manufacture of Al and aluminous refractories.

Bayerite, a-Al(OH)3.

B.C.F., CBrCIF. A gas used in fire extinguishers.

Be. Bervllium.

Beckmann Rearrangement. An intermolecular rearrangement for the conversion of ketoximes to amides, using PCI₅ or SbCl₂ or HoSO4 or HCl.

$$\begin{array}{ccc} C_6H_5-C-C_6H_5 \rightarrow & O=C.C_6H_5\\ \parallel & & \mid\\ N.OH & HN.C_6H_5 \end{array}$$

Beckmann Thermometer. A sensitive mercury thermometer, used for accurate temperature measurement in the determination of molecular weights by boiling point elevation or freezing point depression method. It can be set for any particular range.

Beer's Law. A law stating that the amount of light absorbed depends on the thickness (d) of the absorbing layer and on the molecular concentration (c) of the absorbing medium.

 $I = I_0 e^{-k_0 d}$, where I_0 is the intensity of the incident radiation, and I that of transmitted radiation, and k is molecular absorption co-efficient.

Beeswax. A mixture of mercyl palmitate, cerotic acid and esters, and some paraffins produced by bees. Used in shoe polishes and floor waxes.

Beilstein's Test. A method for detecting the presence of halogens in an organic compound.

Benedict Solution. An aqueous solution of Na2CO3, CuSO4 and sodium citrate used for testing sugars (or other reducing agents). Forms red-yellow or precipitates with sugars.

Benfield Process. A process for removing CO2 from fuel gases.

Benzaldehyde, C7H6O, C6H6CHO. A colourless liquid with alamond like smell, B.P. 180°C. Reduces Fehling's solution and Tollen's reagent. Used in the manufacture of perfumes.

Benzamide, C₆H₅CONH₂. Colourless crystals, M.P. 130°C, B.P. 288°C. A derivative of benzoic acid.

Renzene, C6H6,



An aromatic compound containing

6 C atoms arranged in the form of a regular hexagon, the C-C

bond length is 1 39 Å. Highly inflammable liquid burning with a smoky flame. An important organic solvent, miscible with ethanol, ethanoic acid, ether and propanone. M.P. 5 49°C, B.P. 80°C, D¹⁵ 0.885. Contains 6π electrons represented by the double bonds. Undergoes additions and substitution reactions (electrophilic and substitution reactions). Forms metallic complexes with transition elements. Kekule gave the following structure for benzene:



Benzene Sulphonic Acid, C6H5SO3H. Colourless deliquescent plates,

Benzidine, C₁₂H₁₂N₂. M.P. 127.5°C. Used for blood detections.

Benzoic Acid, C6H5COOH. A white crystalline carboxylic acid, M.P. 122°C, B.P. 249°C, used as a preservative for food and a corrosion inhibitor.

Benzoin C₁₄H₁₂O₂, C₆H₅C(O) CH(OH)C₆H₅. The (±) compound prepared by the action of NaCN on benzaldehyde in dilute

Benzophenone, C13H10O. C6H5C(O)C6H5. Colourless rhombic prisms, M.P. 49°C, B.P. 306°C. Used in perfumery.

Benzoquinone, C₆H₄O₂, (|| ||). Yellow monoclinic prisms,

obtained by oxidising aniline with chromic acid. M.P. 115.6°C.

Benzyl Group. The group C₆H₅CH₂—

Benzyl Alcohol. C₆H₅CH₂OH. Colourless liquid, B.P. 205°C. Used in

Benzyne, C₆H₄. A dehydroaromatic compound. Also known as

- Berkelium, Bk. A radioactive transuranic element of the actinoid series of elements. Does not occur in nature. At. No. 97, M.P. 986°C, ²⁴⁹Bk obtained by the action of neutrons on ²⁴³Am.
- Beryl, 3BeO.Al₂O₃. 6SiO₂. A mineral for the manufacture of Be. Beryl when coloured green by Cr³⁺ are known as *emerald* and when blue green as aquamarine.
- Berrylium, Be. At. No. 4, At. wt. 9'0122, M.P. 1278°C, B.P. 2970°C, D 1'85. The lightest alkaline earth. Grey in colour and is used in nuclear reactors, ceramics, alloys and moderator.
- Bessemer Process. A process for converting pig iron to steel by oxidising the impurities such as, C, Si, P and Mn by air.
- Beta Particle, Beta Ray (β-particles). An electron emitted during some radioactive disintegrations.
- BHC, Benzene Hexachloride, HCH, Hexachlorocyclohexane, $C_6H_6C_6$. An insecticide occurring in many isomeric forms. γ -BHC is also known as *lindane*.

BHT, 2, 6-Di-ter-butyl 4 methylphenol.

Bi. Bismuth.

Bicarbonates. Hydrogen carbonates of various metals.

Bile Acids. Cholic acid.

Bile Salts. The sodium salts of glycocholic acid and taurocholic acids, present in the bile of animals.

Bimolecular Reactions. A chemical reaction involving two molecules, e.g., $2HI \rightarrow H_2 + I_2$.

Binary Compounds. A chemical compound formed from two elements, e.g., KCl or ZnO.

Bioassay. An estimating test (quantitative and qualitative) for substances having a large physiological effect.

Biose. The only carbohydrate with 2 carbon atoms, e.g., glycollic aldehyde (CHO.CH₂OO).

Biosynthesis. A process by which living organisms build up the structures required for their further growth and reproduction.

Bismark Brown. Basic azodyestuff used as hair dye.

Bismuth, Bi. A bright pinkish metallic element belonging to group V of the periodic table. Occurs as Bi₂S₃ and Bi₂O₃. M.P. 271°C, B.P. 1560°C.

Bittern. The liquid left out after sodium chloride has been crystallised out of sea water.

Bisulphan, C₆H₁₄O₆S₂. A cytotoxic alkylating agent used for the treatment of lukaemia.

- Bitumen. A mixture of solid or semi-solid hydrocarbons obtained from coal, oil etc.
- Biuret Reaction (Biuret Test). Substances containing two C(O)NHgroups attached to one another or, to the same N or C atoms, produce a pink or violet colour with NaOH and copper sulphate. Used as a test for biuret, peptides and proteins.
- Bivalent. Showing a valency of two.
- Blast Furnace. A furnace for producing iron from iron (III) oxide.
- Bleaching Powder. A white powder obtained from Cl2 and Ca(CH)2. (slaked lime). Produces Cl₂ on treatment with dilute acids and power is decided by "available chlorine". Used for bleaching paper pulps, for sterilizing water. $Ca(ClO)_2$. $Ca(OH) + CaCl_2 + 2CO_2 \rightarrow 2CaCO_3 + CaCl_2 + 2HClO$.
- Blende. Sulphide ore, ZnS.
- Blue Vitriol (CuSO₄.5H₂O).
- Blue Water Gas. A mixture of CO(41%), H₂(50%) and remainder being N2 and CO2 (is equal amounts) together with a trace of
- BMC. A powerful perchlorinating agent.
- Body Centred Cubic Crystal. A crystal in which the unit cell has an atom, ion or molecule at each corner of the cube and at the
- Boiling Point. The temperature at which the vapour pressure of a liquid is equal to that of the atmospheric pressure.
- Boiling Point-Composition Diagrams. A graph depicting the equilibrium compositions of liquid vapour phases plotted against temperature (at constant P) for a two component system.
- Boltzmann Constant k. A constant, equal to R (Gas constant) divi
 - $k=1.38054 \text{ JK}^{-1}=1.381\times10^{-16} \text{ erg degree}^{-1} \text{ molecule}^{-1}$.
- Bomb Calorimeter. An apparatus for experimental determination
- Bond. A linkage between atoms in molecules and between molecules-
- Bond Energy. The energy involved in forming a bond, e.g., the bond energy for a C—C bond is 348 kJ mole⁻¹. The bond dissociation energy is the energy required to break a particular bond.
- Bonding Orbitals. See molecular orbitals.
- Bone Ash. A white powder obtained by calcining bones (tricalcium

Bone Black, Animal Charcoal. A residue left after the destructive distillation of bones. Used in sugar manufacture as a decolourising agent.

Borane. BH3.

Borates. Salts of boric acid H_3BO_3 , e.g., sodium borate Na_3BO_3 , peroxyborates (used in washing powder) and metaborates $(B_3O_6)^{3-}$ etc.

Borax, Na₂[B₄O₅(OH)₄], 8H₂O. Occurs in nature as kernite (Na₂B₄O₅(OH)₄2H₂O) and tincal (Na₅B₄O₅(OH)₄.8H₂O) and used as a mild antiseptic and a source of boron compounds.

Borax Bead Test. A preliminary test in qualitative analysis.

Boric Acid, H₃BO₃. A white crystalline solid soluble in water and a very weak acid in aqueous solution. Used as an antiseptic, as a food preservative and in the manufacture of pyrex glass.

Born-Haber Cycle. A thermodynamic cycle used to calculate the lattice energies of ionic solids and average bond energies of covalent compounds, e.g., NaCl.

Boron, B. At. No. 5, At. wt. 10.8, M.P. 2300°C. A hard brittle metalloid element of II group of periodic table, occurs in available forms as borates (rasorite, borax and colemanite). Used in nuclear reactors (10B), as borosilicates in enamels and glass and for making light weight components.

Boron Hydrides. Six hydrides of B varying from B₂H₅ to B₁₀H₂₄ containing B-H, B-B and B-H-B linkages.



Used as high energy fuels and a source for pure boron.

Boron Silicates. Silicates containing boron, e.g., danburite (CaB₂Si₂O₈) found in nature and prepared as glasses by fusing together B₂O₈, SiO₂ and a metal oxide (may not be present definitely). Pyrex glasses are borosilicates.

Bosch Process. A method to get H₂ for the Haber's synthesis of ammonia.

Boyles' Law. At a fixed temperature, the pressure of a fixed mass of a gas is inversely proportional to its volume i.e.,

$$pV = K$$
 (Constant)

where K is a constant.

Bromine. Br.

Brady's Reagent. A reagent for testing aldehydes and ketones.

Bragg Equation. An equation to study the crysial structure of a substance (or material) using X-rays. The condition underwhich a crystal will reflect a beam of X-rays with maximum intensity is:

$n\lambda = 2d \sin \theta$

where λ is the wavelength of X-rays, d is the distance between the crystal planes, n is an integer and θ is the angle of incidence (Bragg angle) that the X-ray makes with the crystal planes. Phenomenon is known as Bragg scattering.

Brass. An alloy of Cu and Zn containing up to 30% Zn. Brasses have individual names, depending upon their composition such

Braunite. Brown Mn₂O₃ containing some SiO₂, used as a colouring agent for bricks and pottery. An ore for manganese.

Brazing Metal. See solder.

Breunnerite. Impure magnesite (MgCO₃).

Brine. Concentrated solution of NaCl in water. NaCl is generally

British Thermal Unit (BTU). The amount of heat required to raise 1 lb of water through 1°F under specified condition.

Bromates. Salts containing bromine oxy-anions, hypobromates,

Bromine, Br. At. wt. 79.916, At. No. 35, Adark black liquid, D 3.188, M.P. -7.3°C, B.P. 58.8°C. Red poisonous vapours. Occurs in sea water used in the manufacture of anti-knock agents, flame proofing agents, disinfectants and water purifier.

Bromoform, Tribromoethane, CHBr3. Colourless liquid, M.P. 8°C, B.P. 151°C. Soluble in alcohol and ether.

Bronze. An alloy of Cu and Sn, e.g., Gun metal, beryllium bronze.

Brownian Movement. The random motion of small particles in a fluid, e.g., smoke particles in air. First of all observed with a pollen suspension. The particles large enough to be visible under microscope move freely as they are continuously bombarded by the fluid particles. Used in calculating Avoga-

Brown Ring Test. A test for nitrates (NO₃⁻).

BTX. A mixture of low boiling aromatic hydrocarbons, e.g.,

Buffer. Solutions which resist change in pH. Mixtures of weak acid plus a salt of the acid derived from a strong base, e.t., solution of acetic acid and sodium acetate. The pH of the solution is determined by the dissociation equilibrium of the free acid :

$$e.g., \frac{[H^+] [CH_3COO^-]}{[CH_3COOH]} = K.$$

or by Henderson's equation:

$$pH = pKa + \log \frac{[Salt]}{[Acid]}$$

Buna Rubber. Synthetic elastic substances based on butadiene copolymer. NBR (Buna-M, nitrile rubber) is a copolymer of butadiene (70%) and acrylonitrile (30%) Buna-S is SAR.

Burette. An apparatus used in volumetric analysis.

Burner. A device for the combustion of solid, gaseous or liquid fuels.

Burning Oil. Kerosines oil.

Butadiene, CH₂=CH.CH=CH₂. A colourless gas, B.P. -5°C. Forms complexes, e.g., butadiene-FeCO₃, with metal compounds.

Butadiene Polymer. Polymers derived from butadiene and its deri-

vatives, e.g., styrene.

Butanol, CH3CH2CH2CHO. Colourless liquid with a pungent smell, B.P. 75°C. Used in the preparation of rubber accelerators.

Butane, C₄H₁₀. Lowest member of paraffin series. Exhibits Isomerism, e.g., n-butane (CH₃CH₂ CH₂CH₃) and iso-butane (CH₃CH-CH₃)

Butanoic Acid, C4H8O2. Two acids are known e.g., butanoic acid (CH₃ CH₂ CH₂COOH) and isobutyric acid [(CH₃)₂CHCOOH)].

Butanol, C4O10O. Four butanols are known e.g., normal butyl alcohol (CH₃ CH₂ CH₂ CH₂ OH), isobutyl alcohol [(CH₃)₂ CHCH₂OH], secondary butyl alcohol [(CH₃CH₂CH(CH₃).OH)] and tertiary butyl alcohol [(CH3)3 C.OH].

Butanone, Methyl Ethyl Ketone. C4H8O. CH3 COCH2 CH3. Colourless liquid, B.P. 80°C. Used as a solvent.

Butene, C4O8. Colourless gases with unpleasant odour. Forms three isomers.

Butter of Antimony. SbCl3.

Butyl Rubber. A copolymer of $(CH_3)_2C=CH_2$ (isobutene) and $H_2C=C$ (CH_3) $CH=CH_2$ (isoprene), used as a water proofing membrane. By-product. A substance obtained during the formation of a

chemical product.

C. Carbon.

Ca. Calcium.

Cadmium, Cd. At. No. 48. At. wt. 112'41°C, M.P. 320'9°C, B.P. 765°C, D 8.64. Occurs as Cds in greenockite. Transition metal obtained as a by-product during the extraction of Zn. Used as an anti-corrosion metal, as a neutron absorber in nuclear reactors and also in alkali type batteries.

Cadmium Cell. See Weston cadmium cell.

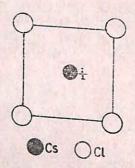
Cadmium Red, Orange Scarlet. A pigment containing CdS (Se).

Cadmium Yellow. A pigment occurring as cadmium sulphide.

Caesium, C. M.P. 28.5°C, B.P. 690°C. D 1.87. At. No. 55, At. wt. 132.90. A reactive element of the alkali metal group. Occurs in silicate minerals, e.g., Pollucite (Cs.Al Si₂O₆). Used in photocells, as a catalyst and in caesium clocks.

Caesium Chloride CsCl. A typical form of structure formed from

Cs2CO3 and HCl.



Caffein, C₈H₁₀N₄O₂. An alkaline present in tea, coffee and guarana. M.P. (anhydrous) 235°C. Acts as a also used in cola drinks, tea and coffee. stimulant and diuretic,

Cage Compounds. Clathrate compounds.

Calamine, ZnCO₃ (British), ZnSiO₄, H₂O (U.S.). Used in lotions,

Calcite. A most stable form of CaCO₃. Have many names, e.g., calcpar, iceland spar, dog tooth spar.

Calcium, Ca. A third element of group II of periodic table, At. No. 20 At. wt. 40.08, M.P. 839°C, B.P. 1484°C, D1.54. Occurs as CaCO₃ (dolomite), CaSO₄ (gypsum), Ca₃(PO₄)₂ halides and silicates (anorthite). Used as a reducing agent in the preparation of metals such as Th, V, Cr and its oxide used in chemical industry. Also used in biological systems. Prepared from CaCO₃ and Al at high temperature.

- Calcium Carbide, Calcium Acetylide, CaC_2 . Manufactured from coke and CaO in the presence of an electric arc. Ionic structure C_2^- and used for the manufacture of cyanamide and ethyne.
- Calcium Carbonate, CaCO₃. A white solid occurs in two crystalline forms, e.g., calcite and aragonite causes temporary hardness in water. Used in Solvay process and for making glass, cement and mortar.
- Calcium Cyanamide, CaNCN. Used as fertilizer.
- Calcium Fluoride, Fluorspar, CaF₂. Occurs as fluorspar. Used in Ca and Al manufacture, in molten salt baths and in optical systems.
- Calcium Hydrogen Carbonate, Ca $(HCO_3)_2$. Formed by the dissolution of $CaCO_3$ by water containing CO_2 . $CaCO_3 + H_2O + CO_2 \rightarrow Ca (HCO_3)_2$

Causes temporary hardness in water.

- Calcium Hydroxide, Ca(OH)₂, (Slaked Lime). A white solid, sparingly soluble in water to form an alkali (lime water). Used as a base, in the manufacture of bleaching powder, mortar and white wash material. Also used for removing temporary hardness of water.
- Calcium Oxide, CaO (Quicklime). A white solid produced by heating Ca in O₂ or by thermal decomposition of CaCO₃. Used in the formation of slag in metal extractions and as a drying agent.
- Calgon. A substance generally, added to detergents to remove unwanted chemicals that have dissolved in water and react with soap to form a scum.
- Caliche. An impure commercial form of sodium nitrate.
- Californium, Cf. At. No. 98. A radioactive transuranic element of actinoid series of metals. The silver-grey metal obtained by neutron bombardment of 243 Am and 244 Cm and 249 Cf.
- Calomel, Hg2Cl2. Mercury chloride.
- Calomel Electrode. A general standard electrode of three types, e.g., 0.1M concentration, 1.0M and saturated KCl. Their respective potentials on the hydrogen scale at 298K are: 0.3338 volts, -0.2800 volts and -0.2415 volts. A half cell having a mercury electrode coated with a paste of Hg₂Cl₂ (calomel) in an electrolyte consisting of saturated KCl solution and mercury chloride solution.

8

- Calor Gas. Calor Propane. The trade name for liquified petroleum gas (LPG) sold in cylinders for domestic use.
- Calorie, Cal. A unit energy approximately equal to 4.2 Joules. The international table calorie is equal to 4.184 Joules (calit). Also used as kilo-calories per gram (for foods) and kilo-calories per kilogram (for fuels) units for the calorific values.
- Calorific Value. The energy liberated by burning unit mass of a fuel (1 ml. for gases).
- Calorimeter. An apparatus for the measurement of calorific values of fuels, foods and for the measurement of heat of chemical reaction.
- Camphor, C₁₀H₁₆O. M.P. 179°C, B.P. 209°C. Obtained from the wood of camphor tree. Used medicinally as a carminative and stimulant, a remedy for colds and in the manufacture of explosives, camphorated oil is 20% solution of olive oil.
- Cane Sugar. Sucrose.
- Cannizzaro Reaction. A reaction producing an acid and alcohol when two molecules of some aldehydes react in the presence of dilute alkalis, e.g., benzaldehyde forms benzoic acid and benzyl
- Capillary Action. A phenomena resulting in the rise or fall of a liquid in a capillary tube when the later is dipped in the liquid.
- Carbanions, R₃C⁻. Ions produced by cleavage of some C-H; C-X; C-metal and C-C bonds. Highly reactive with air
- Carbene, R2C: Contains divalent carbon atom. Produced by pro-
- Carbenium Ions, Carbonium Ions. Positively charged species containing a trivalent carbon atom R₃C⁺. Have strong affinity for nucleophiles, e.g., water.
- Carbides. Derivatives of carbon with elements of lower electronegativity, e.g., B2C, Al4C3. They are extremely hard and infusible and used for making cutting tools.
- Carbinol. Methanol.
- Carbohydrates. A class of compounds occurring in nature and represented by general formula $(CH_2O)_x$ where different values of x represent sugars, starches and cellulose. For exmple in glucose $(C_6H_{12}O_6)$ the value of x is 6 and in sucrose $(C_{12}H_{22}O_{11})$ the value of x is 12. They are classified as: mono, di- or polysaccharides having repeating units equally containing 5 or 6
- Carbolic Acid. Phenol.
- Carbon, C. At. No. 6, At. wt. 12.01, M.P. 3550°C, B.P. 4830°C. Occurs in free as well as combined state.

- Carbon Black. A variety of carbon obtained by the incomplete combustion of natural gas or liquid hydrocarbons. Used in rubber industry, as a decolourizer and as a pigment for paint, ink, etc.
- Carbon Dating. A method of measuring the age of archelogical materials that contain matter of living origin. The radioactive ¹⁴C exchanges (from atmosphere) with ¹²C in living organisms. This exchange stops on death and ¹⁴C decays exponentially. So, the age of once living material can be measured by determining the amount of ¹⁴C present in it.
- Carbon Dioxide, CO₂. A colourless, edourless and non-flammable gas produced by burning of C in O₂. Also produced by respiration. Occurs in traces in atmosphere and liberated by plants. Used as a refrigerant (dry ice or solid CO₂), in fire extinguishers and aerated waters.
- Carbon Disulphide, CS₂. M.P. -112°C, B.P. 46°C. A colourless poisonous liquid. flammable and used mainly as a solvent.
- Carbon Fibres. Graphite fibres used in the strengthening of composites.
- Carbonic Acid, H₂CO₃. A very weak dibasic acid obtained from CO₂ and H₂O.
- Carbonization. To convert an organic compound into carbon by incomplete oxidation at hight temperature.
- Carbon Monoxide, CO. A colourless flammable toxic gas, M.P. -205°C, B.P. -191°C. Formed by incomplete combustion of C or carbon compounds. Strong reducing agent and used in metallurgy.
- Carbon Tetrachloride, CCl₄. Tetrachloromethane. B.P. 78°5C. A colourless non-flammable liquid obtained by the chlorination of methane. Chiefly used as a solvent.
- Carbonyl. A complex in which C=O ligands are co-ordinated to a metal atom, e.g., tetracarbonyl nickel (0), Ni(CO)₄.
- Carboxyl Group. The —COOH group present in organic acids, e.g., CH₃COOH (acetic acid).
- Carboxyl Acids. Organic compounds containing one or more (-COOH) carboxyl group and with the general formula RCOOH. Some of the acids occur in fats and oils and hence the name fatty acids. Named as: mono (containing one -COOH group), di (two -COOH group) and tri (three -COOH) carboxylic acid.
- Carbylamine Reaction. See isocyanide test.
- Carnallite. A mineral of K and Mg, KCl.MgCl₂.6H₂O.
- Carius Method. A process of quantitative analysis for halogens, phosphorus and sulphur in organic analysis.

Caramine. A red lake pigment.

Carnallite, KCl, MgCl2.6H2O.

Carnauba Wax. A yellow, dirty green, brittle and hard wax obtained from the leaves of palm. Used in polishes and varnishes.

Carnolite, KUO2.VO4, 1.5 H2O. An uranium ore.

Carnot Cycle. An hypothetical reversible cycle involving four operations, e.g., Successive adiabatic compression, isothermal expansion, adiabatic expansion and isothermal compression of the working substance. The cycle returns to its initial stage showing that the maximum efficiency for the conversion of heat into work depends only on two temperatures between which the engine works and not on the nature of the material used in the

Caro's Acid, H2SO5. Persulphuric acid.

Carrier Gas. The gas used to carry the sample gas in gas chroma-

Case Hardening. A process for increasing the surface hardness of steel tubes which are used in making gears and crank shafts.

Cast Iron. An alloy of iron and carbon containing about 2.4-4.0% carbon as iron carbide or graphite.

Castner-Kellner Cell. An electrolytic cell used in the manufacture of

Catabolism. A process that breaks down complex molecules to

Catalyst. A substance which when added to a reaction mixture alters the rate of the reaction without itself being changed chemically, e.g., Pt in the synthesis of ammonia. Two types of catalysts: Homogeneous and Heterogeneous.

Catenation. The formation of chains of atoms in molecules.

Cathode. The electrode which carries the negative charge in an electrochemical cell. It is the terminal at which electrons enter the cell. Cations are discharged at cathode.

Cation A positively charged ion, formed by removing electrons from atoms or molecules, e.g., Na⁺, Zn²⁺ or (NH₄)⁺ ions.

Cationic Detergents. Detergents which produce positively charged surface active ions in aqueous solutions. Used in textile industry.

Caustic Potash, 'KOH. Potassium hydroxide.

Caustic Soda, NaOH. Sodium hydroxide.

Cd. Cadmium.

C.D. Circular Dichroism.

Ce. Cerium.

- Celestine, SrSO4. An ore of strontium.
- Cell. A system used as a source of electrical energy. A cell consists of two electrodes.
- Cellophane. A transparent sheet cellulose. Highly inflammable.
- Cellular Plastics. Different varieties of foamed and expanded plastics. Used in packing, thermal insulations and roofing.
- Celluloid. An inflammable thermoplastic.
- Cellulose, $(C_6H_{10}O_5)_x$. A polysaccharide and a chief constituent of the cell walls of plants. Obtained from wood pulp.
- Cellulose Nitrate (gun cotton, nitrocellulose). Highly inflammable substance obtained from cellulose and a mixture of HNO₃ and H₂SO₄. Used as an explosive.
- Celsius Scale. A temperature scale in which the temperature of molten pure ice is taken as 0° and that of boiling water as 100° (at standard P). The degree celsius (°C) is equal to Kelvin.
- Cement. A substance used to make other substances and a powdered mixture of calcium silicates and aluminates obtained by heating limestone (CaO) with clay and grinding.
- Cementite, FeC. A component of some cast irons and steel.
- Centigrade Scale. See celsius scale.
- Centrifuge. A device used to increase the rate of sedimentation of suspensions. Used to separate solid, liquid or two immiscible liquids.
- Centrifugal Pump. A mechanical device commonly used for transporting fluids around a chemical plant.
- Ceramics. Engineering inorganic materials having high melting points. They include silicates and aluminates, refractory metal oxides, metal nitrides, etc., e.g., pottery and porcelain.
- Cerium, Ce. M.P. 799°C. D 6.7. A ductile malleable silvery element of the lanthanide series. Used as a catalyst, in the formation of alloy and glass industry.
- Cerrusite. An ore of lead (PbCO₃) and often found together with galena (PbS).
- C.G.S. System. A unit expressed in the centimeter, gram and the seconds.
- Cf. Californium.
- Chain Reactions. Chemical reactions which proceed by means of a chain reaction mechanism, e.g., the reaction between hydrogen and chlorine:

Cl₂-2Cl⁻ H₂+Cl⁻-HCl+H⁻ H⁻+Cl₂-HCl+Cl⁻

other examples are: Induced nuclear fission reactions (initiated by neutrons).

- Chalcogens. The elements oxygen, sulphur, selenium, tellurium, pollonium of VI group and X² species.
- Chalk. A naturally occurring variety of CaCO₃ formed by marine organisms. Black board chalk is CaSO₄.
- Charcoal. An amorphous form of carbon obtained by heating wood in absence of air. Used as a fuel, reducing agent and adsorbent for gases.
- Charles' Law. At constant pressure the volume of a given mass of a gas is directly proportional to absolute temperature.
- Chelate. A metal coordination complex in which one ligand coordinates at two or more points to the same metal ion, e.g., 1,2—diaminoethane (H₂N CH₂ CH₂NH₂) acts as a chelating agent and beryllium acetyl acetone is a chelate compound.
- Chemical Equivalent The weight of a substance which combines with or displaces 8 parts by weight of oxygen or 35.5 g. of chlorine, etc.
- Chemical Potential (μ) . A measure of the intensity of a chemical substance in a system.
- Chemical Warfare. Materials used in chemical warfare, e.g., toxic chemicals, flame, smoke, defoliating agent, etc.
- Chelmluminescence. The emission of light during a chemical process, e.g., the light emitted by glow worm or oxidation of yellow P.
- Chemisorption. An adsorption of a substance on adsorbate and forming chemical bonds with it.
- Chemotherapy. Selective destruction or prevention of pathogenic
- Ching Clay, Kaolin. A white powder used in paper and pottery industry.

Chinese White. ZnO.

- Chlorinated Rubber. Substances containing up to 70% chlorine obtained by treating rubber with Cl₂. Used as a blending agents for alkylated resins, as an adhesive and film producers in corrosion resistant paints.
- Chlorine, Cl₂. At. No. 17, At. wt. 35.45, B.P. -34.6°C. A gas. Occurs in nature as NaCl, MgCl₂ and in sea water (1%). Obtained commercially by electrolysis of brine and MgCl₂. A bleaching agent and always stored as liquid chlorine (Cl2). Used in the production of anti-knock agents, in water sterilization, pulp and paper manufacture, refrigerants and solvent.
- Chlorine Dioxide, (ClO2). An orange gas used as a powerful oxidising agent.
- Chlorite. A salt of chloric acid.
- Chlorobenzene, C6H5Cl. A colourless liquid obtained from Cl2 and C_6H_6 .
- Chloroform CHCl₃, (trichioromethane). B.P. 60-61°C. A colourless liquid with a sweet pungent smell. Oxidized by air and sunlight to phosgene (decomposition prevented by alcohol). Form iodoform with iodine and KOH. Used as a solvent, volatile anaesthetic.
- Chlorohydrins. Organic compounds containing C (OH). CCl group.
- Chlorophenols. Acidic substances used in phenol resins, anti-septics, disinfectant and germicide.
- Chlorophyll. The green colouring matter of plants. Two chlorophylls are found in plants.
- Chlorophyll a C₅₅H₇₂. MgN₄O₅. (A blue black powder, m.p. 150-153°C), and chlorophyll b, C₅₅H₇₀ MgN₄O₆ (dark green powder, M.P. 120-130°C). It is essential to all life and acts as a catalyst in the photosynthesis of carbohydrates.
- Chloroprene, C₄H₅Cl, (CH₂=CCl. CH=CH₂). Colourless liquid. B.P. 59°C. Used in the manufacture of synthetic rubber.
- Chromatography. A technique of separation of components of a mixture because of the distribution of the components beta mixture because of the solid phase (stationary phase geneween a liquid phase and a solid phase (stationary phase geneween a fiquid phase and which often has a large surface area. Five important types of chromatography are: column chromatography, paper chromatography, thin layer chromatography, gas chromatography and ion exchange.
- Chromium, Cr. At. No. 24, At. wt. 51'9°C M.P. 1900°C. A transition metal occurs naturally as chromite (FeO. Cr₂O₃). Pure tion metal occurs hattanay reduction of $(CrO_4)^{2-}$. Pure chromium obtained by electrolytic reduction of $(CrO_4)^{2-}$. Used

as oxidising agent, for colouring glasses, in textiles, as a catalyst and in leather tanning, etc.

- Chromophore. A group of atoms in a molecule responsible for the colour of the compound, e.g., -C=C-, -C=0, -N=N-.
- Chromyl Chloride, CrO₂Cl₂ A dark red covalent liquid obtained by distilling a mixture of K₂Cr₂O₇ and NaCl with concentrated sulphuric acid. Used as an oxidising agent and as a test for chloride ions in qualitative analysis.
- Chymotrypsins. An important group of proteolytic enzymes which are secreted into the intestines by pancreas.
- Cinchonidine, C₁₉H₂₂N₂O. A stereoisomer of chinchonine.
- Cinnabar, HgS. The chief ore of mercury.
- Cis-. A designation for isomers with groups that are adjacent.
- Cinnamic Acid, C₆H₅CH=CH.COOH. Colourless crystals obtained by Perkin's reaction. Occurs, in sorax or liquid amber. Used in perfumes, flavours, cosmetics, etc.
- Citral, C₁₀H₁₆O. A terpene aldehyde and a chief component of lemon-grass.
- Citric Acid, C₆H₈O₇. A white crystalline tricar boxylic monohydroxy acid found in citrus fruits. Forms three series of salts. Used extensively in the soft drinks and food industries.
- Citric Acid Cycle, Kreb's Cycle. A cyclic process in metabolism which achieves the controlled oxidative breakdown of acetyl co-enzyme A derived from carbohydrates and fatty acids. The products of the cycle are CO₂ and reduced coenzymes (NADH, NADPH, FADH₂, etc.).
- Claisen Condensation. A condensation reaction in which two molecules of ester combine to form a keto-ester in the presence of sodium ethoxide catalyst.

 $2CH_3.CO.O.C_2H_5 \rightarrow CH_3COCH_2COOC_2H_5 + C_2H_5OH$

Clapeyron Clausius Equation. A thermodynamic equation:

$$\frac{d\mathbf{P}}{d\mathbf{\Gamma}} = \frac{\triangle \mathbf{H}v}{\mathbf{T}\triangle \mathbf{V}}$$

where P is the pressure, T is the absolute temperature, $\triangle H_{\nu}$ the molar latent heat of vapourization and $\triangle V$ is the corresponding change in molar volume. The equation is applied to any two phase equilibrium for a pure substance.

- Clark Electrode. See oxygen cathode.
- Clathrate. A molecular compound in which small (guest) molecules are trapped within the lattice of a crystalline (host) compound. Used in the separation of gases.

Claude Process. A process for the liquefaction of air.

Clemmensen Reduction. A reduction leading to the reduction of carbonyl compounds to corresponding hydrocarbons by LiAlH4, etc.

Cleavage. The splitting of a crystal along planes of atoms to form smooth surfaces.

Close Packing. The arrangement of particles in crystals (solids) in which each particle has 12 nearest neighbours in such a way that 6 are in the same layer (or plane) as itself and three each in the layer above and below, e.g., cubic and hexagonal close packings.

Cm. Cerium.

Co. Cobalt.

Coagulation. A process involving the association of particles into clusters, e.g., arsenic sulphide sol can be coagulated by adding unipositive cations.

Coal. A naturally occurring solid fuel existing in the form of seams at various depths below the earth's crust, e.g., peat, lignite. bituminous coal, etc.

Coal Gas. See town gas.

Coalite Process. A process for the carbonisation of coal at 600°C to produce coke, coal-tar, etc.

Coal Tar. A by-product obtained by heating coal in absence of oxygen. A mixture of organic compounds such as benzene, toluene and naphthalene.

Coal Tar Fuels. (C.T.F.). Industrial fuels.

Cobalt, Co. At. No. 27, At. wt. 58.933, M.P. 1475°C. A transition metal, occurs in silver ores (arsenide and sulphides) and Ni, Cu, Pb ores. Used for making alloys and electrical heating elements.

Cobaltite, CoAsS. A cobalt mineral.

Cocaine, C7H21O4. Colourless crystals, occur in coca. A central nervous system stimulant.

Codeine, C₁₈H₂₁NO₃. Colourless crystals, used in the treatment of coughs and as an analgesic.

Code-liver oil. A pale yellow oil extracted from the fresh liver of code.

Codon. See nucleic acid.

Co-enzyme A. An important acyl group transfer co-enzyme in living cells.

- Co-enzymes. Small organic molecules that are necessary for the activity of enzymes e.g., nicotinamide adenine dinucleotide (NAD) and ubiquinone (coenzyme Q).
- Coinage Metals. The metals used to make coins e.g., Cu, Ag, Au.
- Coke. The dense product left in the coke-oven after the carbonization of coal. Used as a fuel, as a reducing agent.
- Coking Coal. The marketable coke obtained after the carbonization of coal.
- Colemanite, CaB₃O₄(OH)₃, H₂O. An important source of borates.
- Collagen. A colourless protein containing chiefly glycine, hydroxy-proteine and proline. Found in all connective tissues of body, e.g., skin, cartilage and tendons.
- Collagenase. An enzyme.
- Colligative Properties. The properties of solutions depending upon the number of particles and not on the nature of particles. Important colligative properties are:
 - 1. The lowering of vapour pressure.
 - 2. The elevation of boiling point.
 - 3. The lowering of freezing point.
 - 4. Osmotic pressure.
- Collman's Reagent. Na₂Fe(CO)₄, 2/3 dioxine.
- Collodion. A nitrocellulose (10.5-12.3% nitrogen), widely used as a base for lacquers, in which a ketone or ester may be used as a solvent.
- Colloid. An heterogeneous system made up of two phases, namely, disperse phase (contain particles, commonly made up of large number of molecules) and the continuous phase (particles are distributed in a continuous medium). A stabilizing agent is used for the stabilization of colloid. For example, Arsenic sulphide sol.
- Colloidal Electrolytes. Soaps and many dyestuffs are colloidal electrolytes.
- Colloidal Mills. Mechanical devices for producing colloidal emulsions or suspensions. Used in pharmaceuticals, paints industry, etc.
- Colorometric Analysis. Quantitative analysis in which the concentration of a coloured solute is measured by the intensity of the colour.
- Colour Developer. Substances used in photographic development
- Colour Indicators. Indicators which depend for their effect on colour change, e.g., acid-base or redox indicators.

- Columbite, (Fe, Mn) (Nb, Ta)2O6. The chief ore of niobium.
- Columbium, Cb. A name of niobium.
- Column chromatography. A technique used for the separation of mixtures of neutral substances with similar physical and chemical properties.
- Combustion. The quick, high temperature oxidation of fuels, converting C to CO₂ and hydrogen to water with the evolution of heat and light.

Common Ion Effect. Consider the following equilibrium reaction:

CD ⇒ C⁺+D⁻

Addition of excess C⁺ or D⁺ to the reaction mixture will shift the equilibrium to the left hand side and thereby, there will be an increase in concentration of CD. This effect is known as common ion effect, used in the qualitative analysis of mixture of salts.

- Common Salt. Commercial name for NaCl.
- Complex. A type of compound in which molecules or ions form coordinate bonds with a metal atom or ion, e.g., [Co(NH₃)₆]Cl₃.
- Complexion. An ion formed by the co-ordination of other ions or molecules to an ion to produce a stable entity, e.g., Fe³⁺ ion and cyanide ions form the [Fe(CN)₆]³⁻ complex ion.
- Complexometric Titrations. Titrations resulting in the formation or decomposition of a complex, e.g., EDTA titrations.
- Component. The number of components in a chemical system is the smallest number of substances which are required to describe the composition of every phase present in the system under study.
- Composites. Plastic matrix reinforced by fibres, e.g., glass, graphite, boron, etc. Used in space crafts, air craft, car components, etc.
- Compound. Substance obtained by combination of atoms of different elements in which the ratio of combining atoms remain fixed and is specific to the substance, e.g., Na₂ CO₃, Ba₃(PO₄)₂
- Concentration. The amount of substance dissolved in a specified weight or volume of the solvent. It can be expressed in various ways, e.g., normality (N), molarity (M), molality (m) and mole fraction (X) of the solution.
- Concentration Cell. A cell which derives its electromotive force from the difference in concentration of solutions of the same electrolyte surrounding the two electrodes.
- *Concrete. The construction material obtained from the hardening of a mixture of cement, sand and water.

- Condensation. The conversion of a gas or vapours into a liquid or solid by cooling.
- Condensation Reactions. A reaction involving the addition of one molecule to another resulting in the elimination of simple molecule such as water, alcohol, ammonia, etc., e.g., Claison
- Condensers. A device used for condensing vapours.
- Conductometric Titrations. A titration in which the end point is determined by observing the change in conductance (conductivity) continuously throughout the addition of titrant and well beyond the equivalence point.
- Conductivity. The reciprocal of the resistance of a circuit (electrolytic solution) and also a property due to which the substance allows the passage of the electric current. The specific conductivity of a substance is the reciprocal of specific resistance. The equivalent conductance (λ) of an electrolyte is given by the

$$\lambda = \frac{1000 \text{K}}{\text{C}}$$

where K=Specific conductance and C is the concentration of

- Condy's Fluid. A disinfectant solution of calcium and potassium
- Configuration. The arrangement of electrons about the nucleus of an atom, e.g., 1s², 2p³, 3d⁵. Also defined as the spatial arrangement of atoms or groups in molecules.
- Conformation. A particular dynamic shape of molecules resulting from the normal rotation of its atoms or groups about single bond, e.g., the two extreme conformation of ethanc are the

Staggered

Conjugate Base. According to Lowry-Bronsted, an acid releases a proton and the ion formed is a conjugate base of the acid.

$$A \rightleftharpoons H^+ + B$$

where A is the acid and B is its conjugate base.

Conjugation. Alternating double (or triple) and single bonds, e.g.,

bonds in
$$-C = C - C = C$$
 or $-C = C - C = O$, etc.

- Conservation of Energy, Law of. In a system of constant mass energy can neither be created nor destroyed but can be transferred into other forms of energy. Energy and mass are interconvertible.
- Conservation of Matter, Law of. The matter can neither be destroyed nor created. There is a conservation of mass and energy together.
- Constantan. A Cu (55%) and Ni (45%) alloy used in resistances and thermocouples.
- Constant Boiling Mixture. A mixture of two liquids constituents in which the composition of vapours is same as that of liquid (azeotropic mixtures).
- Constant Proportions Law of. (Proust's Law). The composition of a pure chemical compound is independent of the method of preparation and that the proportion of each element in a compound is fixed or constant.
- Contact Process. An industrial process for the manufacture of sulphuric acid.

$$SO_2 + O_2 \xrightarrow{V_2O_5} Or Pt.$$

The SO₃ is dissolved in sulphuric acid to form H₂S₂O₇ (Oleum) and which on dilution forms sulphuric acid

$$SO_3 + H_2SO_4 \rightarrow H_2S_2O_7$$

 $H_2S_2O_7 + H_2O \rightarrow 2H_2SO_4$

- Continuity of State. The properties of liquids and vapour become similar at critical temperature. Although the change from liquid to vapour, or vice-versa, is normally discontinuous, a gradual transition is possible with continuity of State.
- Continuous Spectrum. A spectrum composed of a continuous range of absorbed or emitted radiations (well defined lines or bands). A characteristic of unquantized process, such as dissociation.
- Cool Flames. Incomplete combustion shown by hydrocarbons, aldehydes and ethers. Under certain conditions oxidation occurs at low temperatures emitting little heat and light and no flame.
- Cooling Tower. Towers used to cool water used in coolers and
- Co-ordinate Bond (dative bond). A covalent bond formed by sharing of two electrons (lone pair) from one atom by another atom, e.g., the donation of lone pair of ammonia molecule to Cu²⁺ to form [Cu (NH₃)₄]²⁺.
- Co-ordination Number. The number of co-ordinate bonds formed to a metal atom or ion in a complex.

Copolymer. A polymer obtained from the polymerization of two or more monomers, e.g., block and graft copolymers.

Copper, Cu. At. No. 29, At. wt. 63.54, M.P. 1083.4°C. D 8.92. A coinage metal occurring as cuprite Cu₂O, copper pyrites or chalcopyrites CuFeS2, malachile CuCO3. Cu(OH)2 and azurite 2CuCO₃. Cu (OH)₂. A transition metal used in electrical wires and in alloys such as brass and bronze. Forms two series of salts, e.g., cuprous (Cu⁺) and cupric (Cu²⁺) salts.

Copper Bromide. CuBr2 and CuBr.

Copper Chlorides.

Copper (II) Chloride, CuCl2. Dark brown substance, forms complexes such as $[Cu_2Cl_6]^{2-}$ or $[CuCl_4]^{2-}$, etc.

Copper (I) Chloride, CuCl. White solid, forms carbonyl and

Copper Hydroxide, Cu(OH)2. Prepared from copper salts and

Copper Nitrate, Cu (NO₃)₂. A compound forming 9 and 6 hydrates and decomposes on heating to the oxide.

Copper Oxides:

Copper (I) Oxide, Cu2O. A red solid, used in red glasses.

Copper (II) Oxide, CuO. A black solid.

Copper Sulphate.

Copper (II) Sulphate, CuSO4. Blue crystals (blue vitriol) of CuSO₄.SH₂O. The mono hydrate (CuSO₄.H₂O) is formed at 100°C and the anhydrous salt at 250°C. Used in the agriculture

Copper (I) Sulphate, Cu₂SO₄. A grey solid.

Copper Sulphide, CuS. Black solid.

Co-precipitation. The contamination of a precipitate by substances which are generally soluble in the mother liquor is known as

Corrin. Vitamin B12.

Corrosion, A process involving the reaction of a metal with an acid, oxygen, or other compounds resulting in the destruction

Corundum. (emery, Al₂O₃). A naturally occurring form of aluminium oxide that may contain small amounts of iron and silicon oxide, e.g., ruby and sapphire. Used in polishes, abrasives, etc.

Cotton Effect. A phenomena observed in absorption bands.

- Coulomb. SI unit of electrical change, equal to the charge transported by an electric current of one ampere flowing for one second. Faraday correspounds to 96,500 coulombs.
- Coulometer. An instrument used for measuring the amount of electrical charge in a circuit.
- Coulometry. An analytical technique involving the measurement of quantities of electricity.

Coumarin, C9H6O2



Colourless (keto) -

crystals used in perfumery.

- Counters, Radioactive. An instrument for detection and estimation (quantitative) of radioactivity.
- Coupling. A chemical reaction involving two groups or molecules which join together, e.g., the formation of azo compounds.
- Coupling Constant (J). A constant used in nuclear magnetic resonance.
- Covalent Bond. A bond formed by the sharing of an electron pair between two atoms, e.g., a linkage between two hydrogen atoms to form hydrogen molecule.

H·+·H→H··H

Covalent Radius. The equilibrium distance between two atoms joined by a covalent bond of a specific type (single, double or triple) is found to be, with in close limits, constant, e.g., C—C

bond distance in diamond is 1.54 A.

- Cracking. The thermal decomposition of a substance into fractions of lower molecular weight.
- Critical Point. The conditions of temperature and pressure under which the liquid and vapour phases are indistinguishable (both phases co-exist together).
- Critical Pressure (Pc). The minimum pressure required to convert a gas in to liquid at its critical temperature, e.g., critical pressure for carbon dioxide is 73.0 atmosphere.
- Critical Temperature (Tc). The temperature above which a gas can never be liquefied whatsoever high pressure is applied to it, e.g., critical temperature for CO₂ is 31.1°C.
- Critical Volume (Vc). The volume of one mole of a substance at its critical point.

Crotonic Acid

a-Crotonic acid, trans-crotonic acid. Colourless needles, M.P. 72°C

β-Crotonic acid, isocrotonic acid, Cis-crotonic acid. M.P. 14°C,

colourless needles.

Crude Oil. A naturally occurring mixture of hydrocarbons and sulphur, nitrogen and oxygen derivatives.

Crum Brown's Rule. According to the rule a substance C₆H₅A forms m products (disubstituted) if the substance HA can be oxidised directly to HOA; otherwise a mixture of o and p

Crushing and Grinding. A process used in the metallurgical opera-

Cryohydric Point. Eutectic point.

Cryolite, Na₃AlF₆. An important ore of Al.

Cryoscopic Constant. Freezing point depression constant.

Cryoscopy. A method for the determination of molecular weights by freezing-point depression of a solvent.

Crystal. A solid substance bounded by definite faces which intersect at fixed angles (interfacial angles), e.g., cubic, hexago-

Crystal Habit. The description of the way in which the different faces of the same class of crystals grow. Defines the shape of

Crystalline State. Denoting a substance that forms

Crystallite. A small crystal that has a potential to grow larger.

Crystallization. A process for the formation of crystals.

Crystallographic Axes. Imaginary axes defining the repeat unit in a crystal, e.g., axis of rotation of a crystal.

Crystallography. The complete study of the formation, structure

Crystal System. A general classification of crystal systems based on the shapes of their unit cell. Important crystal systems are:

cubic, tetragonal, orthorhombic hexagonal, trigonal, monoclinic and triclinic.

Cubic. Denoting the crystal in which the unit cell is a cube.

Crystal Violet. Methyl violet.

Cs. Caesium.

CS Gas. Malonitriles

CTAB. Cetyltrimethyl ammonium bromide.

Cu. Copper.

Cupellation. A process used in the preparation of Au or Ag.

Cuprammonium, [Cu (NH₃)₄]²⁺. The deep blue copper amine, used in rayon manufacture.

Cuprite, Cu2O. A bright red copper ore.

Curie. The standard unit of radioactivity and equal to the same number of disintegrations as produced by 1 g of Ra $(Ci=3.6\times10^{10} \text{ disintegrations per sec.})$.

Curium, Cm. At. No. 96, M.P. 1340°C. A silvery metal and a potential source of actinides.

Curtius Transformation. A method for obtaining an amine from an ester via the hydrazine, azide and isocyanate.

Cutting Fluids. Fluids used as lubricants and coolants in metal cutting processes.

Cyanic Acid, HNCO.

Cyanine Dyes. Polymethine dyes containing M-C (=C-C)₄=N and relative system.

Cyanoethylation. (A specific Michael reaction). Addition of carbanion species to the double bond of arylonitrile producing a cyanoethyl derivative.

Cyanoferrates. [Fe (CN)₅]⁸⁻ hexacyanoferrates-(III) and [Fe (CN)₆]⁴⁻ and hexacyanoferrates (II).

Cyanogen, (C2N2). A toxic flammable gas.

Cyanohydrins. Organic compounds formed by an addition reaction between an aldehyde or ketone and hydrogen cyanide. Represented by general formula RCH (OH) CN(from an aldehyde) and RR'C(OH)(CN) (from a ketone).

Cyclenes. Cyclic hydrocarbons possessing one or more double bonds.

Cyclic Compounds. A compound made up of a ring of atoms, e.g., homocyclic compounds (those having similar type of

atoms) and heterocyclic compounds (those having different atoms).

- Cycle Oil. A product obtained during cracking process in petroleum industry.
- Cyclic Process. A process in which a system undergoes a series of changes and finally returns to its original state.
- Cyclized Rubber. A modified variety of rubber. Used in corrosionresistant paints and adhesives.
- Cyclobutadiene, C4H4. Highly reactive unsaturated compound and undergoes Diels-Alder reaction.
- Cyclobutane, C_4H_8 . $\begin{pmatrix} CH_2-CH_2 \\ | & | \\ CH_2-CH \end{pmatrix}$. A colourless gas insoluble in water and soluble in organic solvents.

Cyclohexane, C6H12.

A colourless liquid M.P.

6.5°C, B.P. 81°C. Used as an intermediate in the preparation of nylon, solvent for oils and as a paint remover.

Cyclopentadiene, C₅H₆. A colourless liquid, B.P. 42°C insoluble in water, soluble in organic solvent.

and plastics.

Used for the preparation of insecticides

Cyclophanes. Benzene derivatives, e.g., [8] paracyclophane.

Cyclopropane, Trimethylene, C₃H₆. A powerful gaseous anaesthetic.

H,CCCH.

Cyclotron. A device used to accelerate the particles, e.g., protons, by passing them repeatedly through the same electric field.

Cymrose, C7H12O4. A desoxy sugar with a -CH2 group.

Cysteine, C₃H₇O₂N₅. A reduced product of cystine. Present in the body proteins and gets oxidised to cystine on standing in air.

Cystine, Dicysteine, C₆H₁₂O₄N₂S₂. Present in abundance in the proteins of skeletal and connective tissues of animals and in hair and wool.

Cystase. An enzyme which hydrolyses skeletal carbohydrates.

Cytidine, C₉H₁₃O₅N₂. A nucleoside, M.P. 230°C.

Cytochrome. A widely distributed respiratory catalyst associated with the oxidative process of living cell.

Cytoxic Agents. Chemicals injurious to living cells.

D

D. Deuterium.

Dakins Solution. A solution of sodium hypochlorite with 0.5% available chlorine and neutralized with boric acid. Used as an antiseptic for wounds.

Dalton's Atomic Theory. A theory based on the following postulates:

- 1. All elements are made up of atoms.
- 2. All atoms of the same element are identical.
- 3. Atoms can neither be created nor destroyed.
- 4. Atoms combine to form molecules in simple ratios.

Dalton's Law (of partial pressure). The pressure of a mixture of gases is the sum of the partial pressures of each gas present in the mixture. The law is strictly followed by ideal gases.

 $P=p_1+p_2$

where p_1 and p_2 are partial pressures of two gases respectively.

Daniell Cell. A primary cell represented as:

Zn | Zn2+ || Cu2+ | Cu.

where the positive electrode is copper immersed in copper sulphate solution and the negative electrode is zinc immersed in either dilute sulphuric acid or zinc sulphate solution. The e.m.f. of the cell is 1·10 volt.

d-block Elements. The transition elements of I, II and III long periods, i.e., Sc to Zn, Y to Cd and La to Hg. Represented by general formula

 $(n-1)d^n ns^2$, where n=1-10.

DCO. Dehydrated castor oil.

- DDT, Dichlorodiphenyltrichloroethane, C₁₄H₉Cl₁₅. A powerful insectiside, non-phytotoxic to plants and used for controlling mosquitoes.
- **Deactivation.** The process leading to diminish or removal of the chemical reactivity of a substance, e.g., catalyst poisons decrease the activity of Pt catalyst.
- Deaeration. A process for the removal of oxygen and other dissolved gases from solvents by physical and chemical methods.
- de-Broglie Equation. Lous de Broglie (1924) proposed that the wavelength of electrons (λ) is given by the equation:

$$\lambda = (h/p) = (h/mv)$$

where h is the planck's constant, m is the mass of electron, v is the velocity and p is the momentum of electron.

- Debye, D. A unit of electric dipole moment equal to 3.33564×10^{-30} coulomb meter.
- Debye-Hickle Theory. For dilute solutions, Debye-Huckle suggested the following equation:

$$\mu \gamma_i = -\frac{e^3 Z_i^2}{(\epsilon k T)^{1/2}} \sqrt{\frac{2\pi L I}{1000}}$$

where γ_i =activity co-efficient of the ion, Z_i the ionic charge, e=the electronic charge, ϵ =the dielectric constant of the solution, L=Avogadro's number and I=the ionic strength of the solution.

- Decahydrate. A crystalline solid containing ten molecules of water of crystallization per molecule of the compound.
- Decantation. The process involving the removal of a liquid (supernatant) from a suspension or a precipitate.
- Decay Constant. A constant used in a radioactive decay law: $N=No e^{-\lambda t}$

where the number of atoms present at the zero time, N=the number of atoms present at time t and λ =the decay constant $(0.693/t_{\frac{1}{2}})$, $t_{1/2}$ is the half life time.

- Decomposition Voltage. The smallest voltage which is required for the electrolysis of an electrolyte.
- De-emulsification. The process by which emulsions can be broken, e.g., by electrical methods, by chemical methods or by mechanical methods.
- Defect. An irregularity in the lattice of a crystal. Two important
 - 1. Schottky defects. Vacant site with the migrated atom at the surface.
 - 2. Frenkel defects. Vacant sites with an interstitial atom.

- Degenerate Orbital. Specifying different quantum states that have the same energy. In a transition element gaseous atom the five d orbitals are degenerate
- Degree of Hydrolysis (h). The fraction of total salt hydrolysed by water and may be expressed as a percentage.
- Degree of Freedom. From the stand point of statistical mechanics, a degree of freedom means the independent ways in which particle can take up energy, e.g., a monatomic gas has three translational degrees of freedom. From the standpoint of phase, it is the minimum number of variables (e.g., pressure, temperature and concentration) which must be fixed to define the complete state of a system.
- Dehumidification. Removal of condensed water vapours from a vapour gas mixture by condensation.
- Dehydration. Removal of water from a substance.
- Dehydrogenation. A process by which the hydrogen content of a molecule is decreased and the degree of unsaturation is increased, e.g., cyclohexane to benzene.
- Deliquescence. A property of salt which absorbs moisture from the atmosphere and proceeds to dissolve in the absorbed water. The vapour prassure of water over the solid is less than that of the solid surrounding it. The salt is known as "deliquescent".
- Delocalization. Description for molecules or ions in which a bonding electron can not be associated only with one atom but should be considered delocalized over the whole group, e.g., 6π electrons are delocalized over the six C atoms in C_6H_6 .

Delphinin. A pigment

Delphinine. An alkaloid.

Delta Bonding. Lateral overlap between two orbitals.

Delta Metal. Muntz metal.

Demulsification. A process by which emulsions of oil and water can be separated into two liquid phases.

Denaturants. Substances which when added to some other substances make them unfit for human use, e.g., denatured alcohol.

Dendritic Growth. Growth of crystals in a branching habit.

Densitometer. An apparatus for measuring the intensities of lines on a photographic plate, in X-ray analysis and spectrometric analysis.

Density (ρ) or D. The mass per unit volume of a given substance. Units: g. per c.c. or g. per dm⁻³, etc.

- Density Gradient Column. A graduated glass tube, filled with a mixture of solutions to develop a density gradient throughout the length of the tube.
- Depolarizer. A substance used in a volatic cell to prevent polarization such as manganese oxide.
- Depression of Freezing Point. A colligative property or a solution in which the freezing point of the given solvent is lowered by the addition of a solute and is proportional to the number of solute particles in the solution.

Derris. A fish poison.

Desiccant. A material used for drying.

- Desiccator. An apparatus for keeping solids free from moisture or for drying solids.
- Destructive Distillation. A process involving the heating of an organic substance in the absence of air to produce volatile products, which are subsequently condensed and leaving a solid or viscous liquid in the still.
- Desulphorization. A process involving the removal of sulphur compounds from petroleum fractions in the presence of a catalyst. (Also known as hydrotreating or hydrofining.)
- Detergent Oil. Lubricating oil containing detergents. Used in internal combustion engines.
- Detergents. Cleaning agents whose action depends on their power of affecting surface tension and interfacial tension. They wet a variety of surfaces, remove greasy and oily deposits and retain substances. They are water soluble
- Detonating Gas. A mixture of H₂ and O₂ (2:1) obtained by electrolysis of water and explodes to reform H₂O when ignited.
- petrol in a petrol engine, ceases to be practically noiseless and or knocking.

 Describing a process in which the combustion of a sharp metallic hammering is heat. Also known as pinking
- Deuterium, D. At. No. 1, At. wt. 2013. An isotope of hydrogen having diffesent physical and chemical properties. D₂O (heavy preferential electrolysis of H₂O. Used as solvent and as a tracer in chemical and biological processes.
- Deuteron, ²₁D. The nucleus of deuterium atom.
- Devarda's Alloy. An alloy of Al (45%), copper (50%) and zinc (5%). Used in the detection and analysis of nitrates.

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- Developer. Solutions which reduce exposed silver halide grains to silver.
- Devitrification. A process in which under certain conditions the crystallization of glass sets in and the glass becomes opaque.
- 'Dewar' Benzene, C₆H₆. A valence isomer of benzene, to which it reverts on heating.
- Dewar Flask (vacuum flask). A double walled flask made up of thin glass and the space between the walls is evacuated and sealed. Generally, the glass is silvered to reduce radiation.
- Dewaxine. A process used to remove wax from lubricating oil stock.
- Dew Point. The temperature at which a mixture of air and water vapour is saturated with respect to water vapour.
- Dextran. A polymer of glucose made by some bacteria and manufactured by the fermentation of sucrose. Used as anticoagulants.
- Dextrins. Intermediate products formed during the hydrolysis of starch to sugars. Used as adhesives.
- Dextrese, C₁H₁,O₆. Naturally occurring glucese, dextrorotatory, (+) or D⁻(+1). A reducing sugar forming pentacetate osazone, etc.
- Diacetylenes. Hydrocarbons containing two acetylene (C-C≡C-) linkages in the molecules. Mobile colourless liquids having properties similar to ethyne.
- Diagonal Relationship. A relationship which gives rise to similarities in chemical properties (e.g., formation of compounds or formation of complex ions, etc.), between compounds of elements of the main groups of the periodic table, related to one another diagonally. For example: Li-Mg; Be-Al; B-Si.
- Dialysis. A process involving the purification of colloidal sols, e.g., the function of kidneys in mammals is to dialyse the blood.
- Dimagnetism. Describing a property of substances that causes repulsion of the substance from a magnetic field.
- 1-6-Diaminohexane. [H₂N. (CH₂)₆ NH₂]. An organic colourless solid used as a starting material in the production of nylon. Manufactured from cyclohexane.
- Diamond. A hardest, naturally occurring isotope of carbon in which each carbon atom is surrounded by four equally spaced carbon atoms arranged tetrahedrally. Forms a three dimensional net work with a C-C bond length equal to 1.54 A° (equal to that of saturated hydrocarbon) and an angle of 109.5° with

its neighbours. D 3 520 and burns in air or O2 at 700°C to form CO₂ leaving scarcely an ash. Found as basic igneous rock (kimberlite and alluvials). Diamond may be colourless or faintly coloured but is definitely transparent. It has high refractive index and dispersive powers. Used for making "cutting tools". Artificial diamonds are produced from graphite under ligh pressure and temperature.

Diamorphine, Heroin, C₁₂H₂₃NO₅. Soluble in water and alcohol.

Diaphragm Cell. A cell used for the electrolysis of brine.

Diaspore, a-AlO (OH). An important constituent of bauxite.

Diatomic. Describing a molecule containing two atoms, e.g. hydro-

Diazo Dyestuffs. AzO dyestuffs containing -N₂-groups.

Diazomethane, N2CH2. A yellow, poisonous and highly explosive

Diazonium Compounds. An important type of compounds containing the RN=NX group, where R is an aromatic group and

Diazotization. The reaction of an aromatic amine with nitrous

 $C_6H_5 N H_2 + HNO_2 \rightarrow C_6H_5N^+N + OH^- + H_2O$

Dibasic Acid. An acid containing two replaceable hydrogen atoms and forms two series of saits, e.g., H₂SO₄ and H₂C₂O₄ (oxalic

Diborane, B₂H₆. An important hydride of boron. (Also see boron

1,2-Dibromoethane, Ethylene Dibromide. A colourless liquid, B.P. 132°C, M.P. 10°C. Used extensively in petrols and as a

Dicarboxylic Acids. Organic acids possessing two carboxyl (-COOH) groups, e.g., oxalic acid (H₂CO₄). Generally prepared by the oxidation of a glycol, hydroxy-acid or by hydrolysis of cyanoacids. Form both acid and neutral salts.

Dichlorine Oxide, Cl2O (chlorine monoxide). An orange gas formed by passing Cl₂ over mercury oxide. A strong oxidising

Dichroism. Pleochroism in uniaxial crystals.

Dichromate. A salt containing Cr₂O₇²⁻ ion, e.g., potassium dichro÷ mate (K₂Cr₂O₇). They are strong oxidizing agents.

Dielectric Constant. The force F between two electric charges e which are separated by a distance r is given by the equation :

$$F=e^2/Dr$$

where D is the dielectric constant of the medium and is a measure of the polarity of the medium.

Diels-Alder Reaction. See Diene synthesis.

Diene. Organic compounds containing two carbon-carbon double bonds.

Diene Reaction, Diels-Alder Reaction. The 1,4-addition of an alkene or alkyne across a conjugate diene.

Diesel Oil or Diesel Fuel. A petroleum product used as a fuel in diesel engines (commonly known as Derv).

Dieterici's Equation. A modified equation of van der Waal's equation of state.

 $p(v-b) = RTe^{-a/RT}$

where a and b are characteristic constants of a gas.

Diethyl Ether. See ethoxyethane.

Differential Manometer. A manometer used for the measurement of differences of pressure, e.g.. between the vapour pressure of a solvent and its solution.

Differential Scanning Colorimetry. See thermal analysis.

Differential Thermal Analysis. See thermal analysis.

Titration involving the analysis of a sample containing two or more similar reacting species by use of Differential Titration. different reactants or indicators.

Diffraction Pattern. The diffraction pattern formed on a photographic plate when a beam of X rays or electrons is passed through a crystal. Used to study the crystal structures.

Diffusion. Describing the movement of a gas or liquid due to the random thermal motion of its particles. Graham showed that random thermal motion of the particular showed that the rate of diffusion for various gases through a diphragm is its neighbours. D 3.520 and burns in air or O₂ at 700°C to form CO₂ leaving scarcely an ash. Found as basic igneous rock (kimberlite and alluvials). Diamond may be colourless or faintly coloured but is definitely transparent. It has high "cutting index and dispersive powers. Used for making graphite under ligh pressure and temperature.

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- Diazotization. The reaction of an aromatic amine with nitrous acid at low temperatures (below 4°C).
 - $\substack{ C_6H_5\ N \\ Anilne } H_2 + HNO_2 {\to} C_6H_5N^+N + OH^- + H_2O$
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$$\frac{r_1}{r_2} = \sqrt{\frac{d_2}{d_1}}$$

where r_1 and r_2 are the rate of diffusion of two gases and d_1 and d_2 are their respective densities.

Diffusion Pump. A mechanical device to produce high vaccum.

Diffuoromethane CH₂F₂. B.P. -52°C. A gas used as an ingradient in refrigeration.

Digestion. A process involving the recrystallization of a precipitate to make it suitable for easy filtration.

Digol, O.(CH2CH2OH)2. Diethyleneglycol.

Digoxin, C41H64O14. A glycoside used in treating heart failures.

Dihedral Angle. An angle co-relating two parts of a molecule, e.g., in H₂O₂, the angle is 94°.

Dihydrate. A crystalline compound with two molecules of water of crystallization per molecule.

Dihydric Alcohols. See Glycols.

Dihydroxy Acetone, HOCH₂ C(O) CH₂ OH. A strong reducing agent, used to treat sun-tan.

- 1,2-Dihydroxybenzene, Catechol, C₆H₆O₂. Colourless crystals strong reducing. Used as a developer, antioxidant and in the preparation of dyes.
- 1,3-Dihydroxy benzene. (Resorcinol).
- 1,2-Dihydroxyethane, Ethylene Glycol. HO. CH₂. CH₂OH. B.P. 197°C. A colourless, sweet taste and hygroscopic liquid. Used in antifreezes and coolants for engines (50%), in the manufacture of synthetic fibres and as a plasticizers.
- 1,2 Dihydroxypropane, Propylene Glycol. CH₂ CHOH. CH₂OH. A colourless liquid, B.P. 187°C. Used as an anti-freeze and in the manufacture of perfumes.
- Diketones. Organic compounds having two keto (>C=O) groups. Show acidic properties and exist in keto and enol forms.
- Dilatometer. An apparatus used for measuring small changes of volume of a solution, liquid or solid immersed in a liquid.
- Dilute Solution. A solution containing a small amount of solute as compared to that of a solvent.

- Dimer. A compound (or molecule) formed by combination or association of two molecules, e.g., aluminium chloride (AlCl.) is a dimer in vapour (Al₂Cl₆).
- Dimethyl Benzene CaH4 (CH3)2 Xylene. An organic compound hydrocarbon present in the light oil fraction of crude oil. Used as a solvent
- Dimethyl Glyoxime, (CH₃-C=NOH) M.P. 234°C. Colourless

needles, soluble in alcohol. Prepared by the action of hydroxylamine on diacetyl. Forms a dark red crystalline nickel salt. Under suitable conditions it can be used for the detection and estimation of Bi, Cu, Co, and Pd.

- Dimethyl Sulphide, CH₃SCH₃. A colourless liquid, B.P. 37°C. Used for detecting pipe leakage by mixing with gas stream. Also forms complexes with transition metals.
- Dimethyl Sulphoxide, DMSO, Me₂SO. A colourless solid, M.P. 18°C, B.P. 189°C. Used as an extensive solvent and in biology and medicines.
- Dimorphism. Describing the existence of a substance in two crystalline forms.
- 4,6-Dinitro-o-cresol, DNOC, C7H6N2O5. A yellow solid with insecticidal and herbicidal properties.
- Dinitrogen, N2. Describing the N2 group in complexes, e.g., [Ru (NH₃)₅N₂] Cl₂.
- Dinitrogen Oxide, N2O (Nitrous Oxide). A colourless gas with a faint sweet odour, soluble in ethanol and prepared in the laboratory by careful heating of ammonium nitrate: $NH_4 NO_3(s) \rightarrow N_2O(g) + 2H_2O(g)$

Forms N₂ and O₂ on heating above 500°C Used as a mild anaesthetic and produces feeling of elation (laughing gas).

Dinitrogen Tetraoxide, N2O4. A colourless gas, becomes liquid below 21°C and solidifies below -11°C. Forms NO₂ on heating. $N_2O_4(g) = 2NO_2(s)$.

Used as a solvent (liquid) and nitrating agent. 2,4-Dinitrophenyl Hydrazine, CoHoN4O4 (Brady's Reagent). An orange red solid commonly used in solution to identify the a ldehydes and ketones.

Diol (dihydric alcohol), Glycol. An alcohol containing two (-OH) groups per molecule of compound.

- DNBP, C10H12N2O2. Yellow solid, powerful insecticide and herbicide.
- Dobereiner's Triads. Triads of chemically similar elements in which the central element, when placed in order of increasing atomic weight, has an atomic weight approximately equal to the average of the other two, e.g., the triads are recognized as consecutive member of a group of the periodic table (Cl, B, I).
- Dolomite, CaCO₃, MgCO₃. An ore of magnesium.
- Donor. The atom, ion or molecule that donates a pair of electrons in forming a covalent bond.
- Doping. Describing the change in electrical, magnetic and other properties of a solid due to the presence of impurities in the host lattice, e.g., presence of B in Si forms useful conductors.
- d-orbital. Atomic orbital of auxiliary quantum number two containing maximum 10 electrons.
- Double Bond. A covalent bond formed when atoms share two pairs of electrons (double bond), e.g., propene (CH₃CH₂=CH₂).
- Double Salt. Salts formed by mixing equivalent quantities of some salts in aqueous solution followed by crystallization, e.g., Iron (II) ammonium sulphate, FeSO₄ (NH₄)₂ SO₄, 6H₂O.
- Doublet. A term used in NMR (nuclear magnetic resistance)
- Dow Process. A process used to extract Mg from sea water.
- Downs Process. A process for manufacturing Cl₂ by the electrolysis
- Drier, Dryer. A device for drying a material. (Also metallic soaps,
- Drikold. Solid CO2.
- Drilling Fluid. Fluid used to cool the drill, e.g. dirty water.
- Dropping Mercury Electrode, DME. An electrode made up of a column of mercury passing through a fine capillary and emerging in the solution as drops to form a continuous fresh surface.
- Dry Cell. A voltaic cell, chiefly used for flash lights, e.g., the
- Dry Cleaning. A process to clean fabrics using solvents which dissolve dirt at low temperatures, e.g., CaCl4.
- Dry Ice. Solid CO2. Used as a refrigerant.
- Drying Oils. Liquids oxidised by air to dry and hard resins, e.g., linseed oil. Used in coatings, paints, enamels, varnishes, etc.

Dry Point. The temperature at which the last drop of a distilling liquid evaporates.

DSC. Differential scanning calorimetry.

DTA. Differential thermal analysis.

Dulong and Petit's Law. The product of the atomic weight and the specific heat of a metal is a constant (approximately 6.2).

At. wt. \times Sp. heat=6.2

Duma's Method. A method for determining the vapour densities of liquids. Also a method for estimating nitrogen content of a compound by oxidising the compound with CuO followed by reduction of combined nitrogen to nitrogen gas.

Duralumin. A light weight Al alloy containing 3-4% copper with traces of Mg, Mn and Si. Used in aircraft bodies.

Duriron. An iron alloy containing Si (14%), Mn (2%), C (1%), S. (0.1%). Acid resistant. Used in chemical plants.

Dutch Metal. A Zn-Cu alloy.

Dye. A colouring material for fabric, leather, etc. Classified by their ways of application to material, e.g., acid, basic, direct, disperse azo, sulphur and vat dyes.

Dynamite. A mixture of nitroglycerine with other substances to have specific explosive properties.

Dysprosium, Dy. At. No. 66, At. wt. 162.5, M.P. 1409, D 8.5. A lanthanide, used to measure neutron fluxes. Dry compounds are used in lasers and phosphors:



Earth. See alkaline earth metals.

Ebonite, Hard Rubber, Vulcanite. A hard, black, inert material formed by vulcanizing mixtures in which the rubber: sulphur ratio generally lies between 65:35 and 70:30 and less than 4% sulphur remains uncombined.

Ebullioscopic Constant. See elevation of boiling point.

Ebullioscopy. A process involving the determination of molecular weight from elevation of boiling point of solution.

Ebullition. The boiling of bubbling of a liquid.

Ecdysone, C27H44O6. An insect hormone.

Eclipsed Conformation. See conformation.

Edeleanu Process. A process used in petroleum refining.

Edison Cell. See nickel-iron accumulators.

EDTA, Ethlyenediaminetetra-acetic Acid. A compound with mole-

(HOOC. CH₂)₂ N (CH₂)₂ N (CH₂COOH)₂.

Used in the formation of chelates of transition metals.

EFA. Essential fatty acids.

Effective Atomic Number Rule. It is generally found that the total number of electrons available to the central atom from its own electrons by donation (lone pairs of electrons form π bonds), by covalent bonding and taking into account charge add up to the total number of electrons of the next noble gas. Applicable to transition metal carbonyls and many organometallic derivatives but not to many paramagnetic species.

Efficiency (η) . The ratio of a useful energy produced by a system or device to the energy input. For a reversible heat engine the

 $\gamma = (T_1 - T_2)/T_1$

where T₁ and T₂ are the temperatures of source and sink respec-

Efflorescence. The process in which a crystalline hydrated solid loses water of crystallisation to the air.

Effusion. A process involving the passage of a gas through an orifice which has a diameter smaller than the mean free path of the gas molecules. The rate of effusion is proportional to area of the orifice and inversely proportional to the square roots of the

Einstein. A unit used to express radiant energy. The total energy absorbed by molecules in a gram molecule of a substance is Lhv, where L is the Avogadro's number, h is Planck's constant and v is the frequency of light.

Einstenium, Es. At. No. 29.

Einstein's Law of Photochemical Equivalence. Einstein suggested that in a photochemical reaction, each reacting molecule must be excited by absorption of one quantum of light.

Ejector. A device for pumping of gases and liquids.

Elastase. An enzyme.

- Elastin. A protein present in elastic tissues, ligaments and arterial walls.
- Elastomers. Synthetic materials having rubber like properties.
- Electrical Double Layer. An electrical double layer developed at the interface between two phases. One phase attains a positive change and the other phase acquires a negative charge.
- Electrochemical Equivalent, (Z). The mass of an element deposited from a solution of its ions when a current of 1 ampere flows for I second through the electrolytic solution.
- Electrochemical Series. A series describing the activities of metals for reactions that involve ions in solutions.
- Electrochemistry. Describing the branch of chemistry involving the electrolysis and other phenomena when an electric current is passed through an electrolytic solution.
- Electrode. An electro-conducting body that emits or collects electrons or other charge carrier when placed in to an electrolyte.
- Electrodeposition. A process of deposition of a layer of solid (metal) on an electrode by electrolysis.
- Electrode Potential. A measure of the tendency of an element to form ions in solution. It is always defined by comparison with a standard half cell, (e.g., standard hydrogen electrode).
- Electrodialysis. A process by which the electrolytes present in a colloidal solution are removed by dialysis by applying electric filled.
- Electrodispersion. A process for making colloidal solutions when an arc is struck between two metal electrodes under the surface of a liquid.
- Electrokinetic Potential, Zeta Potential. The potential difference across the diffused part of the double layer.
- Electrolysis. The process involving a chemical change (decomposition of a substance) by passing an electric current through an electrolytic solution.
- Electrolyte. A substance which dissociates into ions in solution and hence allows the electric current to flow through the solution.
- Electrolytic Oxidation. A process of oxidation effected by electrolysis.
- Electrolytic Reduction. A process of reduction effected by electrolysis.
- Electrolytic Refining. A method of purifying metals by electrolysis.
- Electromagnetic Radiation. Energy propagated by vibrating electric and magnetic field.

- Electrometer. An instrument used for detecting and measuring the magnitude of an electric charge.
- Electrometric Titrations. Analytical method used in titrations by observing the electromotive force of an inert electrode dipped in solution. Used in titrating coloured solutions.
- Electron. A fundamental particle of negative charge (-1.602192) C and mass of $\frac{1}{1837}$ that of a proton.
- Electronaffinity (A). The energy released when an atom, molecule of group gains an electron in the gas phase to form a negative ions i.e..

 $A(g)+e \longrightarrow A^{-}(g)+energy.$

- Electron Density. A term generally used to explain the probability of finding an electron at a particular point in bonding theories.
- Electron Diffraction. A technique in which a beam of electrons is allowed to interact with atoms or molecules resulting in their Used to determine the structure of substances, mainly the shapes of molecules in gas phase.
- Electronegativity. A measure of the tendency of an atom in a stable molecule to attract electron to itself. In periodic table, elements the left hand side are strongly electronegative than those on hydrogen.
- Electron Exchange. A process of electron fransfer without chemical change.
- Electronic Configuration. Describing a process of distribution of electrons in various shells and sub-shells (orbit and orbitals) of an atom of an element, e.g., the electronic configuration of an element having atomic number 11 is 1s²,2s²2p³3s¹.
- Electron Pair (lone pair). A pair of electrons in an orbital, with
- Electron Spin. A property describing the spin of an electron and having values, $\pm \frac{1}{2}$.
- Electron Spin Resonance. The spatial orientations of the electron spin is aligned either parallel or anti-parallel to the direction to the applied magnetic field.
- Electron Transfer Reaction. Reactions involving transfer of electrons from one group to another, e.g., reduction and oxidation
- Electron Volt (eV). A unit of energy equal to 1.6021917×10⁻¹⁹ Joules. Defined as the energy required to move an electron charge across a potential difference of one volt.

- Electrophile (Electrophillic Reagents). An electron deficient ion or molecule that acts by acquiring electrons, or a share in electrons from other (foreign) molecule, e.g., H⁺, NO₂⁺ ions (positive ions) or SO₃ and O₃ (a molecule that can accept an electron pair).
- Electro-osmosis. If a sol is enclosed by compact diaphragms, so that the motion of the particles be mechanically prevented and the electric current is passed through the two electrodes inserted in the dispersion medium outside the diaphragm, then the dispersion medium moves through the diaphragm towards one of the electrodes. This relative motion of the dispersion medium in the electric field is known as electro-osmosis or endosomsis.
- Electrophilic Substitution Addition. A reaction involving the addition of a small molecule to an unsaturated organic compound, across the atoms held by a double or triple bond, e.g., addition of HBr to ethene:

$$H_2C=CH_2+H^+\rightarrow CH_3CH_2^++Br^-\rightarrow H_3C.CH_2Br$$

Electrophilic Substitution. A reaction involving substitution of an atom or group of atoms in organic compound with an electrophilic reagent, e.g., the nitration of benzene:

 $C_6H_6+NO_2^+\to C_6H_5NO_2+H^+$

- Electrophoresis. The migration of charged particles, colloidat particles or ions through a solution under the influence of an electric field. Used in ionography, zone electrophoresis, electrochromatography, etc.
- Electroplating. The process involving the coating of a solid surface with a metal layer by the application of electrolysis.
- Electrostatic Precipitators. A device (plant) used to remove fine suspended matter from a gas.
- Electrothermic Process. A process used for the preparation of cast steel and other alloys of iron with other metals such as Ni, Cr, etc.
- Electrovalent Bond (ionic bond). A bond formed by the transfer of an electron from one atom to another, e.g., a bond formed in NaCl.
- Electrovalent Compounds. Compounds formed by electrovalent bonding. They are distinguished from covalent compounds by their low volatility, solubility in polar solvents and conductivity.
- Element. A substance which cannot be divided to produce other simpler substances by ordinary chemical methods, e.g., sodium (Na), potassium (K) and aluminium (Al).
- Elementary Particles. Fundamental particles found in nature; e.g., proton, neutron, electron, positron.

- Elements of Symmetry. The symmetry elements of a crystal which describe the space group of a crystal (shape of a crystal), e.g., plane of symmetry, axis of symmetry, point of symmetry.
- Elevation of Boiling Point. A colligative property of solutions in which the boiling point of a solution is raised by the addition of a non-volatile solute. The elevation is directly proportional to the number of solute particles.

 $\wedge T = k_b C_m$

where k_b is a proportionately constant also known as the ebulios copic constant, $\triangle T$ is the elevation in boiling point and C_m is the molal concentration. The units of kb are kelvin kilograms

Elution. The process involving the removal of an absorbed substance in a chromatography column or ion exchange column using a solvent eluent.)

Emarld. The grass green variety of beryl containing some Cr.

Emarld Green, Paris Green. Basic copper ethanoate arsenates (III).

Emery. Impure a-Al₂O₃ (corundum) containing some iron oxide. Used as a polishing agent and abrasive.

Emetin. An alkaloid.

Emission Spectroscopy. A technique involving the excitement of atoms in flames or electrically atoms in flames or electric arc examination of the emission

Emulsion. A dispersed system in which both phases are liquids, generally, one of the liquids is water. Emulsions in which an oil is dispersed in water are known oil in water (O/W) emulsions and those in which water is the disperse phase are water in oil (W/O) emulsions. Used in food (W/O) emulsions. Used in food, cosmetics, horticulture and oil bound water paints, etc.

Emulsion Stabilizers. Weakly absorbed compounds on the grain surface of a photographic emulsion, thereby, displacing the chemical sensitives which in tenulsion, thereby, displacing chemical sensitizer which in turn prevents the formation of fog.

Enamels, Vitreous Enamels. A glassy coating formed on metals by fusion at high temperatures which acts as a decorative and pro-

Enantiomers. Describing a compound whose structure cannot be superimposed on its mirror in of superimposed on its mirror image, e.g., one of any pair of

Enantiotropy. A process in which one allotropic modification of an element or substance is reversibly transformed into another allotrope at a definite transition temperature, e.g., Sn (grey) and Sn (white) have transition temperature 19°C.

- Enclosure Compounds. A prefix used to describe the orientation of atoms or molecules w.r.t. the rest of molecule.
- Endothermic Reactions. Reactions in which heat is absorbed $(\triangle H = +ve)$.
- End Point (equivalence point). The stage in a volumetric titration at which the indicator undergoes maximum change (e.g., colour) for a small volume of added titrant. Alternatively, it detects the end point or equivalence point.
- Energy Levels. Every electronic orbital of an atom is associated with a specific energy value, or energy level. A vibrating or rotating molecule can have discrete vibrational and rotational energy levels.
- Energy Profile. A diagram representing the changes in the energy of a system during the course of a reaction.
- Enols, Enolic Compounds. Organic compounds containing the α -CH group and >C=C-OH grouping (one in which a hydroxyl group is attached to the carbon of a double bond). A tautomeric form of some ketones.
- Enthalpy (H). A thermodynamic state function and a sum of the internal energy (U) and the product of pressure (P) and volume (V) of a system:

H=U+PV

At constant pressure, $\triangle H = \triangle U + P \triangle V$.

Entropy (δ). For a reversible system, the change of entropy is defined as the heat absorbed divided by thermodynamic temperature:

 $\delta S = \delta Q/T$

A thermodynamic quantity and a measure of the disorder of a system. The higher the entropy, the greater is disorder, e.g., $S_{gas} > S_{liquid} > S_{solid}.$

- Enzymes. Proteins that catalyse a specific biochemical reaction. Present in all living organisms and catalyse most of the cell reactions, e.g., the hydrolysis of fats, sugars, proteins and their resynthesis. Obtained from plants, animals and micro-organisms. They have a high degree of specificity. An increase in temperature decreases the rate of enzyme action. They are destroyed by much change in pH values and inactivated by low conscentration of active metals.
- Epimerism. A variety of isomerism in which isomers differ in the position of -OH groups, e.g., a and β form of glucose (epimers).
- Epoxide. Three membered ring compounds (organic) in which two carbon atoms are bonded to the same oxygen atom, e.g.,

Epoxy. A prefix indicating a group in a molecule, e.g., epoxyethane.

Epoxy Resins. Polyethers produced by condensation of epichlorohydrin with polyols such as bisphenol A, or by epoxidation of Diels-Alder adducts with peroxy compounds. Used as adhesives,

Epsom Salt. MgSO₄.7H₂O.

Equilibrium Constant. In a chemical reaction of the type $aA+bB \rightleftharpoons cC+dD$

the equilibrium constant (k) is defined as $K = [C]^{a}[D]^{a}/[A]^{a}[B]^{b}$

where [A], [B], [C] and [D] represent the active masses of reactants and products respectively and a, b, c and d denote their number of moles. For a gaseous reaction,

 $K_P = [(p_C^o)(p_D^d)]/[(p_A^a)(p_B^b)].$

where pa..... etc. are the partial pressures of A.....etc.

Equilibrium, Metastable. If a system is capable of undergoing a spontaneous change thermodynamically, persists without change, the system is said to be in a metastable state.

Equipartition of Energy. The total energy of a molecule is, on average, equally distributed among the available degrees of

Equivalent Conductivity. The conductivity of an electrolytic solution containing 1 g. equivalent of the electrolyte and is equal to the product of specific conductance and volume (ml) containing 1 g

Equivalent Weight. The number of grams of an element that could combine with or displace one gram of hydrogen (or 8 grams of oxygen or 35.5 grams of chlorine), e.g., equivalent weight of hydrochloric acid is 36.5(35.5+1.0). It is also the atomic weight

Equivalent Proportion Law. Substances (elements and compounds) react together in the mass ratio of their equivalents. Er. Erbium.

Erbium, Er. At. No. 68, At. wt. 167:25, M.P. 1522°C. D. 9:01. A soft malleable silvery element of the lanthanide series of metals. Occurs in association with other lanthanides. Used in nuclear and glass industry.

Erdmann's Salt. NH₄ [Co(NH₃)₂(NO₂)₄].

erg. A unit of energy used in C.G.S. system and equal to 10-7 joules.

Es. Einstenium.

ESCA. Electron spectroscopy for chemical analysis.

ESR. Electron spin resonance.

EFA. Essential fatty acids, e.g., lionoleic acid.

Essential Oils. Volatile oils extracted from plants and used in the purified form as source of compounds.

Esterification. A reaction in which an acid reacts with an alcohol to form an ester and water, e.g.,

 $CH_3COOH + C_2H_5OH \rightleftharpoons CH_3COOC_2H_5 + H_9O.$

- Esters. A type of organic compounds formed by the reaction of acids with alcohols and represented by general formula RCOOR', where R and R' are organic groups, e.g., ethylacetate.
- Ethanal, CH₃CHO, (Acetaldehyde). Colourless liquid with a typical smell, B.P. 20.8°C, miscible with water, alcohol and ether. Insoluble in concentrated calcium chloride solution. Used as a chemical intermediate for the formation of various organic compounds.
- Ethanamide, CH₃CONH₂, Acetamide. Long white needles which absorb water resulting in liquid. M.P. 82°C, odour of mice, weakly basic.
- Ethane, C₂H₆. A colourless, odourless gaseous alkane. B.P. -89°C and used in low-temperature refrigeration plant.
- Ethane-1, 2-diol, CH₂(OH)CH₂(OH) (ethylene glycol, glycol). syrupy organic liquid generally used as anti-freeze and in the manufacture of Terylene.
- Ethanoic Acid, CH₃COOH, (Acetic Acid). Colourless liquid with a pungent irritating odour, B.P. 119°C. Manufactured from oxidation of ethanol at 60°C. Used as ethanoic anhydride for oxidation of ethanol at 60°C. the manufacture of cellulose ethanoate, etc.
- Ethanol, C2H5OH, (Ethyl Alcohol). Colourless volatile liquid, occurs in intoxicating drinks and is obtained by fermentation of sugar:

 $C_6H_{12}O_6 \longrightarrow 2C_2H_5OH + 2CO_2$

Only 15% concentrated alcohol is produced by yeast and higher only 15% concentrated around by distillation. Used as a solvent.

Ethanoyl Chloride, CH3COCl, (Acetyl Chloride). Colourless liquid with a pungent odour, fumes in moist air and used as an acetylating agent.

- Ethene, C₂H₄. (Ethylene). A colourless, etheral odour gas occurring in natural gas crude oil and coal gas. M.P. -169°, B.P. -105°C. Highly reactive, unsaturated compound. Reacts with hydrogen bromide, hydrogen iodide, ammonia, water and oxygen under suitable conditions. Forms plastic polythene, styrene vinyl
- Ethene Polymers, Polyethene, Polythene. Most important polymeric substances, used in fabrication, pipe, where coating, textiles Chlorinated polymers are used for flooring.
- Ether. A type of organic compound containing the group—O—and ethers represented by the general formula R-O-R', where R and R' are alkyl or aryl groups. Used formerly as an anaesthetic
- Ethyl Alcohol. See ethanol.
- Ethyl Amine, C₂H₆NH₂. A colourless liquid amine. Used in the
- Ethylene. See ethene.
- Ethylene Oxide, C_2H_4O , $\begin{pmatrix} CH_2 \\ | \\ CH_2 \end{pmatrix}O$ A colourless gas, used as a fumigant and insecticide.
- Ethyl Ethanoate, (C₂H₅O-O.C.C.H₃). An ester used as a solvent for plastics and in perfumery and flavouring.
- Ethyl Lactate, C₈H₁₀O₃. A colourless liquid, B.P. 154°C, used as a solvent for cellulose nitrate and acetate and for some resins
- Ethyl Silicate, Silicon Ester, Si (OC₂H₅)₄. A thin mobile liquid, B.P. 168°C, obtained by the reaction of alcohol with silicon tetrachloride Decomposed by the reaction of alcohol with silicon tetrachloride. Decomposed by water and used for water proofing,
- Ethyl Vinyl Ether, CH₃ CH₂ OCH=CH₂. B.P. 35°C. Used in the
- Ethyne, C2H2, Acetylene. See acetylene.
- Eu. Europium.
- Eucalyptus Oil. Oil distilled from the leaves of various species of Eucalyptus. Used mainly as an antiseptic and as a prophylactic for cold and influenza.
- Eudiometer. An apparatus used in the volumetric analysis of gases.
- Europium, Eu. At. No. 63, At. wt. 1520. Least abundant rare earth element which is used as a neutron absorber in neutron

- Eutectic. A mechanical mixture of two substances which melts sharply and the temperature at which it melts is known as eutectic temperature.
- Eutectic Point. The point on the phase diagram of a mixture of substances which represents the melting point and composition of an eutectic is known as an eutectic point.
- EVA Plastics. A copolymer (vinyl ethanoate).
- Evaporation. All liquids and solids exhibit a vapour pressure at a temperature. Some molecules in the liquid have sufficient energy to escape into the gas phase resulting in the build up of vapour pressure over the liquid surface. If this process takes place in a closed container, an equilibrium is established between the molecules that leave the liquid surface to form vapours and the molecules which return to the liquid from vapour phase. No such equilibrium is established, if this process occurs in an open container. The liquid molecules are continuously removed from the liquid surface. This phenomena is known as evaporation.
- Evipan. A trade name for hexobarbitone.
- EXAFS. Extended X-ray absorption fine structure spectroscopy.
- Exchange Reaction. A reaction which occurs without a chemical change, e.g., $H_2+D_2\rightleftharpoons 2HD$. Such exchange reactions are the basis for the separation of isotopes.
- Excitation. The process resulting in the excited state of an atom, molecule, etc.
- Excitation Energy. The energy needed to push an atom, molecule, etc. from lower energy state to a higher energy state. Also defined as the difference between two energy levels of the system.
- Excited State. A state of an atom, molecule, etc., having an energy greater than that of the ground state.
- Exclusion Principle, Pauli's. No two electrons in an atom can have all the quantum numbers identical.
- **Exothermic.** Describing a chemical reaction in which heat is evolved $(\triangle H = -ve)$, e.g., combustion.
- Explosion. A process of rapid combustion in homogeneous fuelair mixtures with the flame passing through the mixture from the source of ignition. Generally, it takes place at constant pressure and volume.
- Explosives. Solids, liquids or gases which when subjected to a local impulse, e.g., shock, friction, sparks, etc., undergo rapid decomposition with the formation of a huge quantity of heat and large volumes of gases which may occupy many times the volume of the original explosive, e.g., trinitrotoluene.

External Indicator. An indicator used outside the reaction container, e.g., potassium ferricyanide in iron titrations.

Extinction Coefficient (a). A characteristic quantity of a medium which absorbs light and is the reciprocal of the layer thickness measured in centimeters, in which the intensity of the incident radiation has been reduced to 1/10 of its original value. It may

$$\alpha = 0.4343 \text{ K}$$

where K is the absorption co-efficient.

Extraction. The process involving the removal of a soluble component from finely divided solids by means of a suitable solvent.

Extra Nuclear Electrons. , See electronic configuration.



F. Fluorine.

F Centre. An anionic site in a crystal occupied by an electron.

Face-centred Cubic Crystal, (f.c.c.). A crystal structure in which the unit cell has particles te.g., atoms, ions or molecules) at each corner and each face centre of a cube.

Face-centred Lattice. A lattice which has an atom at the face-

FAD. Flavin adenine dinucleotide.

Fahl Ore, Cu₃Sb₃.

Fahrenheit Scale. A temperature scale having the ice temperature as 32°C and the steam temperature as 212°C. A correlation between degree Celsius (C) and Fahrenheit (F) is given by the

 $\frac{C}{5} = \frac{F-32}{9}$

Fajan's Method. A volumetric method for the titration of Cl with Ag⁺ using fluorescein as an adsorption indicator (end point is the formation of red precipitate).

Fajan's Rule. A rule describing the variations in the degree of covalent character in electrovalent (ionic) compounds in terms of polarization effects. Ionic compounds are readily formed (1)

by an atom, which yields ions with a low charge (e.g., not more than Al3+), (2) by large cations (Cs+ should form more electrovalent link than Na), and (3) by small anions (Cl more than I-).

'Farad (F). The SI unit for capacitance. A Farad is the capacitance developed when the plates of a capacitor are charged by one coulomb and the applied potential difference is 1 volt.

1F=1 coulomb volt.

Faraday (F). A unit of electric charge equal to 9.648670×104 coulombs.

Faraday Effect. The phenomena in which a plane polarized light is rotated when passed through a solid or liquid placed in a magnetic field.

Faraday's Laws of Electrolysis. The law states :

- (1) The amount of chemical change occurring at an electrode is strictly proportional to the quantity of electricity passed through the solution.
- (2) When different compounds are decomposed by the same quantity of electricity, the amounts of products obtained at different electrodes are proportional to their chemical equivalents.

Esters of fatty acids with glycerol having general formula: Fats.

CH₀.OOCR₁ CH.OOCR, CH₀.OOCR₃

where R1, R2 and R3 may be the same fatty acid residue. But, where R₁, R₂ and are mixed glycerides, each fatty acid being in general, fats are mixed glycerides, each fatty acid being different. They are the essential constituents of animals.

Fatty Acids. Monobasic acids containing carbon, hydrogen, oxygen and an alkyl radical attached to the carboxyl group. oxygen and an analysis acids are represented by general formula. The saturated latty acid, palmitic acid, etc.), and the Various unsaturated acids are represented by general formulae various unsaturated actos are represented by general formulae $C_nH_{2^{n-2}}$, (e.g., acrylic acid), $C_nH_{2^{n-4}}O_2$ (e.g., linoleic acid series) and $C_nH_{2^{n-6}}O_2$, (e.g., linolenic acid series). Occur in nature mainly as glycerides and as esters of alcohols.

f.c.c. Face-centred cubic lattice.

Fe. Iron.

Fehling's Solution. A solution used to test reducing sugars and the -CHO group (aldehyde group). A solution of copper the -CHO group (aldehyde group). the -CHO group (and any as gloup). It solution of copsulphate, sodium potassium tartarate and sodium hydroxide.

- Feld Scrubber. Bartlett-Harber washer.
- Feldspar. A group of aluminosilicates containing one or more of the followish bases: potash, soda, lime, barium oxide. Used in the ceramic and enamelling industries.
- Femto, (f). 1 femtometer (fm)=10⁻¹⁵ meters (m).
- Fenton's Reagent. An aqueous solution of Fe2+ salt, (e.g., FeSO₄) and H₂O₂ used in oxidising polyhydric alcohols.
- Fermentation. A chemical reaction brought about by microsorganisms (moulds, yeast or bacteria), e.g., the formation of ethylalcohol from sugars.
- Fermi. A unit of length equal to 10⁻¹⁵ metre.
- Fermium, (Fm). A radioactive transuranic element of actinide series. Does not occur in nature.
- Ferrates. Oxyanions of iron, e.g, [FeO₄]²⁻, Sr₂FeO₄ and Na₄[Fe(OH)₆], etc.
- Ferredoxin. Non-haem iron containing proteins.
- Ferricyanides. Salts of hydroferricyanic acid, e.g., potassium ferricyanide K₃Fe(CN₆). Used in quantitative analysis and in the laboratory as a test for ferrous iron.
- Ferrites. Compounds of ferric oxide with a basic oxide. Used in electronic industry.
- Ferritin. A soluble protein responsible for absorption and storage of iron in the body.
- Ferrocyanides. Salts of the hydroferrocyanic acid H₄[Fe(CN)₆], e.g. potassium ferrocyanide (K₄[Fe(CN)₆], H₂O).
- Ferromagnetism. Describing magnetic behaviour showing large magnetic susceptibility and the magnetic moment does not fall to zero when the applied magnetic field is removed.
- Ferromanganese. A type of pig iron containing 6% to 7% carbon and enough of Mn. Used for recarburization in steel manufacture.
- Fertilizers. Substances of N, P or K, etc., which when added to soil increase production, e.g., urea, ammonium sulphate, ammonium nitrate, etc.
- Fibres. Materials used in textiles, e.g., wool, cotton (natural fibres), asbestos, glass wool (inorganic fibres) and synthetic fibres (polyester, nylon).
- Fibring en. A protein present in animals.
- Fibringon. A protein present in silk fibre.

Fieser Solution. A solution used to remove O₂ from, e.g., N₂. Anaqueous alkaline solution of sodium anthraquinone β-sulphonate reduced with sodium diathionate.

Fillers. A solid material such as slate fibre, glass fibre, mica, cotton wool, etc., which are used to improve the physical properties or reduce the cost of production of synthetic compounds, (e.g., rubber, plastics, etc.).

Film. A material deposited in thin layer form.

Filter. A device used for filtering process.

Filter Aids. Material used to increase the rate of filtration, e.g., kieseguhr.

Filter Press. A mechanical device used for filtration under pressure.

Filter Pump. A type of vacuum pump used in the laboratory for vacuum filtration, distillation and creation of low grade vaccum.

Filtration. Describing a process used to remove suspended particles from a fluid (liquid) by passing the liquid through a porous material (the filter paper).

Fine Structure. Closely spaced lines observed at high resolution in a spectral line or band.

Fire Clay. A refractory clay used for the manufacture of furnace bricks, crucible, etc., and contains chiefly aluminium silicates.

Fire Extinguishers. Materials used to extinguish or prevent propagation of flame, e.g., water, NaHCO₃ solution, CO₂ foam formed from a carbonate and an acid, liquids such as CCl₄ and solids such as sodium bicarbonate and potassium bicarbonate.

Fire Point. The temperature at which a substance, when ignited, first burns freely when the ignition agent is removed.

Fire Retardant. A material used to resist burning.

First Order Reaction. The reaction in which the rate of the reaction is proportional to the concentration of one of the reacting species. Alternately, the time required for the completion of species. Alternately, the time required for the completion of any fraction of this reaction is independent of initial concentration of reactant. The first order rate equation is given by

$$k = \frac{2.303}{t} \log \left(\frac{a}{a - x} \right)$$

where k is first order rate constant, a is the initial concentration of reactant (molar) and (a-x) is the concentration of reactant at time 't'.

- Fischer-Topsch Reaction. The catalytic hydrogenation of hydrogen and carbon monoxide, watergas, at 200°C to produce hydrocarbons of high molecular weight.
- Fission Products. The products of the fission of heavy nuclei and generally very radioactive.
- Fittig Reaction. See Wurtz reaction.
- Fixation. Describing a process used to remove unwanted silver halide from a protographic material by fixing material (sodium thiosulphate) after development. Important fixers are: thiocyanates, cyanides, thiourea, etc.
- Flame Emission Spectroscopy. Emission spectroscopy in which the emitting atom is present in a flame.
- Flame Front. The region between the luminous zone and the dark zone of unburnt gases. Used in the measurement of burning velocity of gases.
- Flame Retardants. Materials which when added to fabric, reduce tendency to flame and burn by glowing, e.g., borates, phosphates, bromine, Sb₂O₃, etc.
- Flame Test. A preliminary test applied in qualitative analysis to identify the elements present in a sample. The colour of the flame indicates the presence of an element in the sample, e.g. barium salts impart green colour to bunsen flame.
- Flare Stack. A chimney at the top of which unwanted gases are burnt.
- Flash Back. Describing the flame speed of the combustion mixture in the burner. To maintain a stationary flame, it is necessary to balance the flame speed of the combustion mixture and the velocity of the mixture in the burner tube.
- Flash Distillation. Describe a process of rapid removal of solvent.

 Used in distillation of sea water.
- Flash Photolysis. A technique for investigating free radicals in gases. A powerful flash of light of very high energy is passed through for a few micro seconds resulting in the decomposition. Measureusing an oscilloscope, and the kinetics of very fast reactions can thus be investigated.
- Flash Point. The lowest temperature at which enough vapour is given off by a flammable liquid to ignite in the presence of a catalyst.

Flavone C₁₅H₁₀O₂. A group of yellow pigments, occurs naturally

as dust on the flowers and leaves of primulas, can be synthesised from o-hydroxyacetophenone and benzaldehyde.

Flavones. A group of yellow pigments, occur in the plant kingdom. related to flavone and found in association with tannins.

Flint. A compact mass of quartz. Used in the manufacture of porcelain.

Flocculation. Describing the process of destruction of colloidal sol. A term often used with coagulation and precipitation.

Florisil. Magnesium silicates which are used in chromatography.

Floatation. A process used for the concentration of ores (e.g., sulphide ores). It involves the formation of a moderately stable foam by addition of a small quantity of a suitable reagent (e.g., pine oil) followed by aeration. The particles of ore collect in the liquid air interface of the bubbles and foam is continuously removed from the top of the cell.

Floatation Cells. A device used for the concentration of ores.

Fluidization. Describing a process involving the suspension of mass of a solid particles in an upward-flowing stream of gas or liquid.

Fluon. A variety of polymers.

Fluorene, C₁₃H₁₀. Colourless flakes exhibiting a violet fluorescence.

Fluorescein, C20H12O4. Red crystals with green iridescence M.P. 315°C. Shows intense green fluorescence in alkaline solutions. Used as a dyestuff and as a detector for water leaakage.

Fluorescence Indicator. An indicator which is used in the titration

of deeply coloured or turbid solutions.

Fluoridation. The addition of small quantities of fluoride compounds into the water supply to minimise the tooth decay in

Fluorides. The salts of hydrofluoric acid. Fluorimetry. An analytical technique using the fluorescent pro-

perties of the substances to be estimated.

- Fluorine, F. At. No. 9, At. wt. 18'9984, BP. -188'14°C. The lightest, greenish-yellow and most reactive gas of halogen series. Chief ore is fluorite (CaF2). Manufactured by the electrolysis of molten KF/HF electrolytes, using copper or steel apparatus. Elementary fluorine exist as F₂. Fluorine gas, HF and fluorides are toxic. Its compounds are used in the preparation of inert plastics, refrigerants, aerosol propellants, toothpaste and for
- Fluorine Polymers. Polymers containing fluorine, e.g., Teflon, polyhexafluoropropene, polyvinyl fluoride, etc.
- Fluoroacetates. Derivatives of fluoroacetic acid, CH2FCOOH. Colourless, very stable and highly toxic substances.
- Fluoroapatite, Ca₅(PO₄)₃F. Naturally occurring phosphate rocks.
- Fluoroborates. Salts of fluoroboric acid.
- Fluoroboric Acid, HBF₄. Obtained from BF₃, B(OH)₃ and water. Contain tetrahedral (BF₄)⁻ ion which is weakly basic in nature.
- Fluorocarbons. A class of organic compounds with general formula CxFy, e.g., C₆F₆, The C-F bond is quite stable and long chain molecules are easily formed. Show very similar freezing and boiling points to the corresponding hydrocarbons. Obtained by the fluorination of hydrocarbons with fluorine. Fluorocarbon oils, grease and inert dielectrics are used under those conditions in which ordinary material will be attacked. Chlorofluoro carbons are used as refrigerants, aerosol agents and propellants.
- 1-Fluoro-2, 4-dinitrobenzene, Sanger's Reagent, C6H3FN20. A reagent for the identification of the terminal amino group of
- Fluoroform, CF2H.
- Fluorophosphates. A group of substances with general formula R₂PO₃F, where R is an alkyl group. Highly toxic substances.
- Fluorosilicates. A group of substances containing (SiF₆)²⁻ and
- Fluorosulphuric Acid. HSO3F.
- Fluorspar. A naturally occurring form of CaF₂. Used in certain
- Flux. A substance used to prevent oxidation (no oxide on metal

An additive substance used in smelting metals to react with silicates and other impurities to form a low melting slag, e.g., CaCO₃.

Fluxional Molecule. Describing a molecule which arranges so easily at room temperature that the normal concept of structure is inadequate i.e., the relative position (of atoms of molecules) become indistinguishable. No structure exists for longer than 10⁻² seconds. Example PF₅ (a trigonal bipyramid) exhibits only one type of fluorine ¹⁹F by n.m.r. spectrocopy.

Fly Ash, Na₂CO₃. Sodium carbonate.

Fm. Fermium.

FMN. Flavin mononucleotide.

Foams. A coarse dispersion of a gas in a liquid. Most materials capable of lowering surface tension considerably (e.g., soaps) lead to foaming. Foaming is developed by agitation of liquids and gases in the presence of stabilizing agents. Foams are used in rubber preparation and in fire fighting.

Folic Acid, Vitamin Bo.

Describing several closely related compounds with vitamin activity. Folic acid and its derivatives are widely spread in nature. Green leaves are very rich in this vitamin. Used in the treatment of sprue and some other types of anaemia, and required for the growth of some bacteria.

Food Additives. Substances added to foodstuffs to preserve and improve favour, e.g., calcium phosphate (caking agent), colouring agents, nutrients, preservatives, etc.

Formaldehyde, HCHO. A colourless gas with a characteristic and pungent odour. The commercial solution 'formalin' contains 40% formaldehyde. It acts as a powerful germicide (in solution and vapour form) and hardens tissues. Used for the tanning of leather, preservation of skins, as an antiseptic, for the manufacture of synthesis of resins, plastics and various dyes.

Formalin. See methanol.

- Formic Acid, Methanoic Acid, HCOOH. A colourless fuming liquid, B.P. 100.5°C, miscible with water and alcohol. Occurs in sweat and urine. Used as a food preservative, solvent and as a reducing agent.
- Formol Titration. A method for estimating carboxyl groups in amino acids and free carboxyl group in proteins (Sovensen
- Formula. A representation of a chemical compound using symbols for atoms and subscript number to show number of atoms present in the compound, e.g., sodium dichromate (Na₂Cr₂O₇).
- Formyl. The C(OH) group.
- Formylation. The process of introducing a formyl group in organic
- Fr. Francium
- Fraction. Describing a mixture of liquids with similar boiling. points collected by fractional distillation.
- Fractional Crystallization. A process of crystallization (or separation) of the components of a mixture in a solution. Used for the
- Fractional Distillation. A process of separating a mixture in to aseries of fractions of different boiling points (different volatili-
- Fractionating Column. Rectifying columns used in fractional distilla-
- Francium, Fr. A radioactive element of alkali-metal group occurring in uranium ore in a small quantity. Electronic configuration 7s¹, M.P. 27°C and forms single series of Fr+ salts.
- Franck-Condon Principle. The time needed for an electronic transition in a molecule is very much less than the period of vibration of the constituent nuclei of the molecule. Used in the energy changes and spectra molecules.
 - Frasch Process. A process for obtaining sulphur.
- Free Energy. A measure of the ability of a system to perform useful work. A thermodynamic state function (G). The free energy change $(\triangle G)$ in a system is given by the equation: $\Delta G = \Delta H - T \Delta S$

where $\triangle H$ is the enthalpy change, $\triangle S$ is the change in entropy and T is the thermodynamic temperature. $\Delta G=0$, at equilibrium state. Free energy change is expressed in kilojoules per

- Free Radical. Molecules or ions with unpaired electrons. They are formed by breaking a covalent bond, e.g., CH₃Cl→CH₃·+Cl·. They are highly reactive. The most stable free radicals are NO, NO₂ and molecular oxygen, O₂. They can be stabilised and isolated under special conditions.
- Freezing. Describing the process of converting a liquid into a solid.
- Freezing Mixture. A mixture of substances used to produce low temperatures, e.g., a mixture of sal (NaCl), ice and water can produces a temperature of -21°C. Other freezing mixtures are NH₄Cl/ice (-15°C), alcohol/solid CO₂ (-72°C), etc.
- Freezing Point. The temperature at which a liquid is in equilibrium with its solid phase at a standard pressure. It is equal to the melting point of the solid.
- Freezing Point Depression. The depression produced by dissolving one mole of the solute in one litre of solvent is called the molecular depression or cryoscopic constant.
- Freons. A trade name for a group of halogenated hydrocarbons containing fluorine and chlorine. They are easily liquefied gases and are used in refrigeration and air-conditioning. The important freons are:

12, dichlorodifluoromethane $CHCl_2F(B.P.-33^{\circ}C)$ 21, dichlorofluoromethane $CHCl_2F(B.P.~9^{\circ}C)$.

Frequency, v.- The number of wavelengths passing a fixed observer (or point) per second. Frequency is reciprocal of wave number and proportional to energy.

Freundlich Isotherm. An empirical relation given by the equation

$$\frac{x}{m} = kc^{1/2}n$$

where x is the amount of the substance adsorbed, m is the weight of adsorbent, c is the concentration or pressure and k and n are constants. n is usually 0.2 to 0.1.

Friedel-Crafts Reaction. A method employed for the substitution of an alkyl or acyl group on to a benz-ne ring (using an electrophilic reagent). The acylating or alkylating reagent may be an acid chloride (RCOCI) or alkyl halide (RX), alcohol (ROH), etc:

 $C_6H_6+RCOCI\rightarrow RCOC_6H_5$ $C_6H_6+CH_3I\rightarrow C_6H_5CH_3+HI$

The aromatic hydrocarbon is refluxed with alkylating or acylating agent in the presence of anhydrons, aluminium chloride

catalyst which accepts a lone pair of electrons on the chlorine polarizing the chloroalkane to produce positive charge on the carbon followed by electrophillic substitution. The reaction is useful for the synthesis of hydrocarbons and ketones.

Froth Flotation. A process for the purification of ores.

Fructose, C₆H₁₂O₆. A sugar present in fruit juices, honey and cane sugar. Exists as a ketohexose in a pyranose form when free.

D-Fructose, C₆H₁₂O₆. Crystallizes as large needles, M.P. 102-104°C. Occurs in fruit juice (with glucose). Soluble in water, twice as sweet as glucose and exists in nature as furanose form, but

L-Fucose, C₆H₁₂O₆. Microscopic crystals, M.P. 145°C. Present Furanose form in the blood group polysaccharides and in seaweed.

Fuel Cell. A type of cell, similar to galvanic cell in which fuel is converted directly into electricity. The fuels used may be gaseous or liquid, e.g., hydrogen, hydrazine, methanol, ammonia, etc., and oxidants and generally oxygen or air. Fuel and oxidants are supplied to the respective electrodes which are usually porous and activated by catalyst. Large fuel cells can produce tens of amperes (electromotive force is about 0.9V and efficiency about 60%). Batteries of fuel cells are used in aircrafts.

Fuel Oils. A group of petroleum products (oils) used for the production of heat or power, e.g., engine fuels and burner fuels.

Fuels. Materials used for the production of heat energy, e.g., fuel oils, synthetic gases, rocket fuels, nuclear fuels, etc.

Fuller's Earth. A natural clay used as an adsorbent and industrial

Hydrocarbons represented by general structure given Fulvenes. below

$$\begin{array}{c|c}
H & H \\
3 & 4 \\
\hline
 & 5 \\
2 & 1 \\
H & H
\end{array}$$

$$\begin{array}{c|c}
R \\
R'$$

where R and R' are usually hydrogen, alkyl or aryl-group.

Fumarase. An enzyme used in the formation of citric acid [via (-)malic acidl.

Fumaric Acid, C4H4O4, Trans-butenedioic Acid.

HOOC.CH HC.COOH

Colourless needles, M.P. 300-302°C. Forms maleic anhydride and (±)-malic acid. Used as a food acid.

Functional Group. A group of atoms in an organic compound which is responsible for the characteristic properties of the compounds, e.g., alcohol (-OH), aldehyde (-CHO), ketonic (>C=O), aldehydic (>C=O), amino $(-NH_2)$, nitro $(-NO_2)$, etc.

Fundamental Unit. The units of length mass and time, e.g., metre, kilogram and the second in SI units.

Fungicide. Materials used to prevent fungal attack, e.g., in agriculture, wood, etc. Example: Cu, Sn, S, dithiocarbonate, etc.

Furan, Furfuran, C4H4O. A colourless liquid, B.P. 32°C.

Forms resins in the presence of mineral acids.

Furanose. A form of sugars existing in α and β forms.

Furfural, Furfuraldehyde,

A colourless liquid, darkens an exposure to light. B.P. 162°C. A colourless liquid, darkend fuel oils. Used as a solvent for Occurs in essential oils and fuel oils. Used as a solvent for decolourizing resins and as a solvent.

Furfural Alcohol, $C_5H_6O_2$. Colourless liquid, B.P. 170°C. Poisonous in nature, used in the formation of resins.

Furoic Acid, Pyromucic Acid, C5H4O3,



Colourless crystalline solid, M.P. 125-132°C, B.P. 230-232°C.

Fuse Oil. A mixture of fatty acids, alcohol and esters produced as a high boiling fraction during the distillation of fermentation

Fused Ring. See ring.

Fusidic Acid, C₃₁H₄₈O₆. An antibiotic.

Fusion. Melting.



Ga. Gallium.

Gabriel's Reaction. A reaction involving the conversion of a halogen compound into the corresponding amino compound by treatment with potassium phthalimide followed by hydrolysis. Used

Gadolinium, Gd. M.P. 1313°C. A ductile malleable silvery element of the lanthanide series of periodic table. Used in ferrites, in alloys and electronic industry.

Galactans. Polysaccharides (e.g., agar) which occur in wood and

D-galactose, C₆H₁₂O₆. An isomer of glucose present in plants. A Galena, PbS. A main ore of lead.

Gallates. Anionic gallium containing species.

Gallic Acid, 3,4,5 trihydroxybenzoic Acid, C7H6O5. Colourless crystals with one molecule of water of crystallization. Occurs in wood tissues, gall-nuts, tea and in tannins. Obtained from tannins by hydrolysis.—Used in the manufacture of ink. Forms porogallol. Gallium

At. No. 31, At. wt. 69.72, M.P. 29.8°C, B.P. 2403°C, D 5.907. Occurs in Zn and Al minerals. A soft silvery lowmelting metallic element belonging to group III of periodic table. Used as a semi-conductor (Gallium arsenide), low melting alloys and the high temperature thermometers.

Galvanic Cell. See cell.

Galvanizing. A process for coating steel with Zn for protection by dipping into molten Zn or by electro-deposition (cold galvanizing).

Gamma Radiation. A form of electromagnetic radiation emitted by nuclei of atoms. The Y-rays constituting the radiation are of very short wavelength and very hard X-rays. γ-rays have high penetrating power. The energy of a y-photon is given by the equation

W = hv

where h is Planck constant and v is the frequency of radiation.

Gamma Rays. See Y-radiation.

Gammexane. See BHC.

Gangue. The clay and other silicates occurring in an ore.

Garnets. A group of silicates, M₃" M₂'" (SiO₄)₃ (M"=Ca, Mg, Fe: M"'=Al, Cr, Fe), having discrete SiO₄⁴⁻ groups. Used as gunstones and an abrasive.

Gas. The most diffused state of matter in which forces of attraction between the molecules of the substance are very small (e.g., van der Waals' forces). The molecules move freely and continually collide with each other and with the walls of the container. The pressure of the gas is due to the collision of the molecules with the walls of the container.

An ideal gas follows the gas laws e.g., Boyles' law, Charles' law, Avogadros' law etc. All real gases deviate from the ideal behaviour.

Gas Analysis. A process used to analyse a mixture of gases e.g., by absorbing each constituent by a reagent (e.g., CO₂ in KOH), adsorbing the gas on a substance (e.g., H₂ adsorbed by Pd), vapour phase chromatography, titration as in volumetric analysis (e.g., H2S and NH3), etc.

Gas Calorimeter. An apparatus to measure the calorific value of a

Gas Chromatography. A technique employed for the separation and analysis of mixtures e.g., gas chromatography employes a column packed with either a solid stationary phase (gas-solid chromatography or GSC) or a solid coated with a non-volatile liquid (gas liquid chromategraphy or GLC).

Gas Constant, R. A universal gas constant derived from general gas equation PV=RT. R has the following values:

- 1. 0.082 litre-atmos per degree per mole.
- 2. 8.31432×107 ergs per degree per mole.
- 3. 8:31432 joules per degree per mole.
- 4. 1.987 cal. per degree per mole.

- Gashydrates. Clathrates formed by water with some gases and organic molecules e.g., Cl₂ 7.3 H₂O.
- Gasification. A process by which a solid or liquid feedstock is transferred to a gaseous fuel (lower molecular weight) containing lower carbon to hydrogen ratio e.g., production of town gas, or substitute natural gas (SNG) from petroleum and the produc-

$$C+H_2O \rightleftharpoons CO+H_2$$
water gas

- Gas Laws. The laws which describe the behaviour of gases e.g., Boyles' law, Charles' law, general gas law, etc.
- Gas Oil. The petroleum fraction intermediate between kerosene and light lubricating oil. Used as a diesel fuel.
- Gasoline. A mixture of hydrocarbon fractions used as a fuel for motor and aviation. Aviation gasoline may contain branched
- Gastrin. A hormone of gastro-intestinal tract.
- Gattermann's Reaction. A method for introducing chlorine, bro-
- Gattermann's Synthesis. A method for the synthesis of aromatic hydroxyaldehydes e.g., AlCl₃ is used to carry out the condensation of phenol with a mixture of gaseous hydrochloric acid.

 $C_6H_5OH + HCN + HCl \rightarrow p \cdot C_6H_4(OH)CH = NH.HCl$ JH2O P-C₆H₄(OH)CHO+NH₄CI

- Gauss, G. The unit of magnetic flux density in the C.G.S. system
- Gay-Lussac's Law. Gases combine in volumes that are in simple ratios to each other, and to the products if it is also gaseous. The law holds goods for ideal gases e.g., I volume of nitrogen combines with 1 volume of oxygen to form 2 volumes of nitric
- Gel. Hydrophillic colloids that are usually stable but may be made to coagulate partially under certain conditions, such as lowering the temperature. This produces a pseudo-solid or easily deformable jelly-like mass of intertwining filaments which may enclose the whole of the dispersion medium. Two important 1. elastic gel e.g., gelatin.

 - 2. rigid gel, e.g. silica gel.
- Gelatin Hardness. Substances added to harden gelatine.
- Gelatin. A protein occurring in bones and in fibrous tissues in the form of anhydride, collagen, which on boiling with dilute acids

forms gelatin. Gelatin swells in cold water, but is insoluble in it. Used in cooking and photography.

Gel Filtration. A technique used to separate molecules according to size. It is mainly employed in biochemistry. Used to determine the molecular weight of proteins and in the purification of proteins.

Gelignite. A variety of dynamite.

Gem Position. Describing the position in a molecule on the same atom e.g., 1, 1-dichloroethane is a gem dihalide (CH3CHCla).

Gemstones. Materials used in jewellery e.g., a-alumina (corundum) forms ruby.

Geometrical Isomerism. See Isomerism.

Geon. See Freons.

Germanates. Oxy compounds of germanium e.g., SrGeO3 and Mg.GeO4.

Germanes, Germanium Hydrides. A very pure source of Ge e.g., GeH4, Ge2H6, etc.

Germanium, Ge. M.P. 937'4°C At. No. 32, At. wt. 72'59, D 5'35. A hard brittle grey metalloid, occurs in Zn ores (in ZnS) and a member of group IV of periodic table. Also found in argyrodik (4Ag₂S.GeS₂). Used as an alloying agent, catalyst, phosphorus etc.

Germanium Oxide, GeO₂. Obtained by heating Ge in oxygen. Occurs in two varietes; one of which is soluble in water and

the other is insoluble in water.

Getter. A substance used to remove impurities e.g., Li is used to remove O, S, etc., from Cu and Cu-alloys.

The proportionality factor used in the co-relation of magnetic moment (μ) and the number of unpaired electrons (s)

$$\mu = g\sqrt{s(s+1)}$$
g=2.03 for a free electron.

Gibb's Equation. An equation used for single non-ionic solute in dilute solution

$$\Gamma = -\frac{c}{RT} \frac{d\gamma}{dc}$$

where γ is the surface tension of the solution, Γ is the amount where y is the surface tonsion of the surface, R is the gas constant and c is the concentration per unit volume.

Gibbs Free Energy. See free energy.

Gibbs Helmholtz Equation. For a chemical reaction taking place at a constant pressure

$$\triangle \mathbf{G} = \triangle \mathbf{H} - \mathbf{T} \triangle \mathbf{S}$$

where $\triangle G$ is the Gibbs free energy change, $\triangle H$ is the enthalpy change and $\triangle S$ is the entropy change. For a reaction taking place at constant volume

$$\triangle A = \triangle E + T \left(\frac{dA}{dT}\right)_{v} = \triangle E + T \triangle S$$

where $\triangle A$ is the change in Helmholtz free energy and $\triangle E$ is the change in internal energy.

Used to calculate heats of reaction.

- Gibbsite, 7-Al (OH)3. A variety of aluminium hydrate (monoclinic) and is calcined prior to use.
- Gilsonite. A natural high melting point bitumen (M.P. 170°C). Has a high di-electric strength and very low water absorption.
- A carminative which is used with purgatives to prevent griping. It is warming and stimulating the stomach and is added to medicines as a flavour.
- Girard's Reagent. A reagent used to separate aldehydes or ketones from other neutral compounds. They are quaternary ammonium salts of the type Me₃NCH₂CO NHNH₂+X⁻ which form water soluble salts with carbonyl compounds.
- A hard transparent material obtained by heating calcium oxide (lime), sodium carbonate, and sand (Si (IV) oxide). It is regarded as a super cooled liquid and resistant to chemical attack. In some varieties of glass, silica has been replaced by other substances such as B2O3 or P2O5; etc.

The important varieties of glass are:

Soda glass (calcium silicates), borosilicate glass (containing boron oxide in the glass), flint glass (K2O or BaO as the constituent

Glass Electrode. A thin-walled glass bulb immersed in a solution containing H⁺ (ions) and platinum wire dipping in a solution of an electrolyte which fills the bulb. The glass surface developes a potential with the solution(of H+ ions) and the thin glass serves as a suitable conductor between the outer and the inner

Glass Fibres. Glass filaments added to resins to give them strength.

GLC. Gas liquid chromatography.

Globin. A protein which combines with haem to form haemoglobin. Globulins. A widely distributed class of proteins found in milk

Glove Box. An apparatus used to manipulate moisture or O2

Glucans. Polymers of glucopyranose.

- Gluconic Acid, Dextronoic Acid, C₆H₁₂O₇. Colourless crystals, M.P 165°C, soluble in water and alcohol. Its calcium salt is used in medicines.
- Glucose, C₆H₁₂O₆, Dextrose. An hexose widely present in plants, fruit juices and in the sugar of blood. Obtained from cellulose, starch, sucrose and other polysaccharides by acid or enzyme hydrolysis. It exists in a number of forms.
 - 1. Aldehyde (only in solution)

2. a- and β-Glucopyranose (containing carbon atom 1 as asymetric).

3. α- and β-glucofurcnose (unstable and known only in solution)

Simple glucose is α -glucopyranose monohydrate, M.P. 80.85°C and $[\alpha]p+113.4$ °C.

Glucosides (Glycosides). A sugar derivative e.g., methyl glycoside ($C_0H_{11}O_5-O-CH_3$). They are isomeric as they contain asymetric C atom.

Glucuronic Acid, C₆H₁₀O₇. An oxidant product of glucose, M.P. 165°C. An important constituent of haemicellulose and plant gums.

Glue. A colloidal mixture of proteins, made up from amino-acids (condensation of). It is prepared from waste skin, bones, tendons, etc. 'Fish glue' is prepared from fish refuse.

Glutamic Acid, C5H9NO4, a-Aminoglutaric acid, CH (NH₂) COOH

CH₂

CH₂-COOH

A naturally occurring substance and is dextro-rotatory, [α]ρ²⁵+11.0°. Present in large amounts in protein hydrolysis

Glutaric Acid, Pentanedioic Acid, C5H8O4. M.P. 97-98°C.

Glutathione, Glutamylcysteinylglycine, GSH, C₁₀H₁₇N₃O₆S. 190-192°C. A tripeptide and a constituent of all cells. It acts as an oxygen carrier.

Gluten. A mixture of gliadin and glutelin proteins obtained from

Glyceraldehyde, C₃H₆O₃, OHC. CH (OH). CH₂OH. A colourless

Glyceric Acid, 2, 3 dihydroxypropionic Acid, C₃H₆O₄. An optically

Glycerides. Esters formed by alcohols and carboxylic acids. Classified as mono-, di- and tri-glycerides depending on the number of acid radicals combined with the three hydroxyl

Glycerol, Glycerine, 1, 2, 3-trihydroxypropane, C₃H₈O₃, CH₂OH. CH(OH).CH2OH. A colourless, odourless, viscous liquid with a sweet taste. M.P. 20°C, B.P. 182°C/20 mm. Occurs in all animal and vegetable fats and oils. Commercially obtained as a by-product in soap industry. Forms nitroglycerine with HNO₃, acrolein with H₂SO₄ or KHSO₄. Used as a solvent, as a moistening agent for tobago, in the manufacture of explo-

Glycerol Phosphoric Acid, 3-phosphoglyceric Acid. prepare medicinal salts such as calcium salt of glycerophosphoric Glycin. See amino phenols.

Glycine, Glycocoll, H₂N.CH₂.COOH. M.P. 260°C. A hydrolysis product of amino acids and is the simplest amino acid. It has

Glycogen, $(C_6H_{10}O_5)_x$ (animal starch). A carbohydrate store of animals and is made up of many glucose units linked together. Molecular weight about 4 million. It is readily hydrolysed in the digestive system to glucose (by glucosidases). It is stored mainly Glycol. 1, 2-Dihydroxyethane.

- Glycollic Acid, Hydroxyacetic Acid, CH2OH.COOH. Colourless crystals, M.P. 80°C. Occurs in the juice of sugar cane and beets. Obtained by electrolytic reduction of oxalic acid. Used in textile and leather industry as a cleaning agent.
- Glycols, Dihydric Alcohols. Organic compounds derived from aliphatic hydrocarbons by replacing two H atoms by hydroxyl (-OH) groups, e.g., ethylene glycol (CH₂OH.CH₂OH). All are colourless liquids, miscible with water and alcohol. Obtained by oxidation of olefins by potassium permanganate.
- Glycolysis. A process in which the metabolic breakdown of carbohydrates in living organism, takes place.
- Glycosidases. An enzyme.
- Glycosides. See glucosides.
- Glyoxal, Biformyl, Ethane dial, CH(O).CHO. M.P. 15°C, B.P. 51°C, soluble in water. A yellow solids obtained by the oxidation of acetaldehyde or ethylene by selenium oxide.
- Glyoxylases. An enzyme used in the preparation of lactic acid.
- Glyoxylic Acid, CH(O).COOH. M.P. 98°C. A thick syrup, widely present in plant and animal tissues. Condenses with urea to form allantoin.
- Glyptals. Resins.
- Gold, Au. At. No. 79, At. wt. 196 9665, M.P. 1064 4°C, D 19 3. A transition metal that occurs native, unreactive, very ductile and malleable. A good conductor of heat and electricity. Forms colloidal solution e.g., by reducing gold chlorine solution with hydrazine. It dissolves in aqua regia and halogen solutions. Attacked by fluorine. Used in jewellery, as an i.r. reflector, as a radiation source, etc.
- Gold Chloride, AuCl₃. A compound obtained by dissolving gold in aqua regia. It exists as a dimer (Au₂Cl₆) and is used in photo-
- Gold Grains. A radiation source having encapsulated 198Au in Pt.
- Gold Number. A term used in colloidal sol. Gold number of a protective colloid is the number of milligrams of the dry colloid which when present in 19 ml. of a standard red gold sol (0.0055% Au) will just prevent its change of colour from red to blue on the addition of 1 ml. of a 10% NaCl solution. Smaller the gold number of a substance, more efficient it is for protection
- Goldschmidt Process. A process for extracting some metals from their oxides by reduction with Al.
- Gold, Standard. Pure gold is very soft for coinage or ornaments and is alloyed with Cu or Ag. Pure gold is 24 carat and forms five is alloyed with Cu or Ag. Full gold. Purity is expressed in parts standard alloys, 22, 18, 15, 12 and 9. Purity is expressed in parts per thousand or carats.

Gold Sulphide, Au₂S (grey) and Au₂S₃ (black).

Gonadotropic Hormones. Several peptide hormones effecting the activity of gonads (sex hormones).

Gouy Balance. A balance used to determine magnetic suscepti-

Graham's Law of Diffusion. The rates of diffusion of gases are inversely proportional to the square root of their densities, i.e.

$$\frac{r_1}{r_2} = \sqrt{\frac{d_{\rm B}}{d_{\rm A}}}$$

where r_1 and r_2 are the rates of diffusion. Principal is used in

Graham's Salt, NaPO₃. A polymeric meta phosphate.

Gram Atom. Expressing atomic weight in grams.

Gram Equivalent. The equivalent weight of a substance expressed in grams e.g., one equivalent of sodium hydroxide weighs 40

Gramicidine. An antibiotic.

Gram Molecular Volume. The volume occupied by one gram molecule of an element or compound in the gaseous state. At standard temperature and pressure the gram molecular volume for all elements and substances is 22.414 litres.

Gram Molecule. The weight in grams of a substance.

Granite. An igneous rock. Used as a building stone and road metal.

Granulation. A process of size, reduction and size enlargement for producing granules. Dry granulation forms pellets from dry materials, which are crushed into the required size. In wet granulation, the paste formed with a liquid is dried and cut into Grape Sugar. Dextrose.

Graphite. An allotrope of carbon, a good conductor of heat and electricity and also known as 'black lead'. The atoms are arranged in layers (sheets). The C-C distance between the layer is 3.40A°

Used as a lubricant, refractory, as a moderator and reflector in nuclear reactors, in pencil makings, paints, etc.

Graphite Carbon. Carbon combined with or dissolved in iron or

Gravimetric Analysis. A technique of quantitative analysis invol ing precipitation, drying and the final weighing of a stable derivative.

Gravitation. An elementary particle which exists in wave form in

- Greases. Solids or semí-fluids obtained by intermixing of a thickening agent in a liquid lubricant. Resistant to temperature. Greases may contain soaps of Na, Ca, Al, etc. Non-soap greases contain organo-clays (Benton grease) or some dyestuffs.
- Grignard Reagent. Compounds of the type R-Mg-X obtained by the action of alkyl and aryl halides with Mg in the presence of dry ether. They are highly reactive and react with:
 - (a) alkyl and aryl halides to form hydrocarbons.
 - (b) metal halides to form organometallics.
 - aldehydes and ketones to produce secondary and tertiary alcohols.
 - (d) amides and nitriles to give ketones.
 - (e) water and dilute acids to form hydrocarbons.

Grinding. A process of crushing.

Grottius Draper Law. The law states that the only absorbed light (photons) by a substance can effectively induce a chemical change. All the absorbed light is not used for a chemical change.

Ground State. The lowest energy state of an atom, molecule or ion.

Group. A group in the periodic table is a series of chemically similar elements that have similar electronic configuration e.g., the general electronic configuration of I group elements is represented by outer s¹ configeration.

Guaicol, C₇H₈O₂. M.P. 32°C. A component of guaiacum resin. Used in the preparation of vanillin and denatured alcohol.

Guaioi, C₁₅H₂₆O. M.P. 93°C. An alcohol.

Guanase. An enzyme.

Gum Acacia, Gum Arabic. Obtained from acacia trees and is used in the manufacture of plasters, in food industry, etc.

Gums. A gum obtained from plants e.g., gum acacia, gum traga-canth. Soluble in water and insoluble in organic solvents. A term used in petroleum industry for a 'dark coloured polymer'.

Gun Cotton. See cellulose nitrate.

Gun Metal See bronze.

Gun Powder. A powdered explosive containing sulphur, charcoal and potassium nitrate.

Gutta-percha. A naturally occurring polymer which is used as a rubber additive obtained from tropical trees. Gypsum, Selenite, CaSO4, 2H.O. A mineral used in the manufac-

ture of plaster of paris, as a filler, etc.



h. See hapto.

H. Hydrogen.

Ha. Hahnium.

Haber Process. An industrial process for the manufacture ammonia from N₂ and H₂ in in the presence of a catalyst at low temperature (450°C) and high pressure 250 atmospheres.

 $N_2+3H_2\rightleftharpoons 2NH_3$

The catalyst used can be finally divided iron (traces of Mo or Ca), ferric oxide (containing traces of silica and K₂O) etc.

Haem, C₃₄H₃₂O₄N₄Fe. The non-protein portion of the haemoglobin molecule containing iron in the ferrous, state, obtained by

Haematin, C₃₄H₃₃O₅N₄Fe. M.P. 200°C. A blue black crystalline solid obtained from haemin. It resembles haem and contains iron in

Haematite, Fe₂O₃. A mineral and chief ore of iron. It is isomorphic with corundum and has blood red colour in massive formation.

alkaline solution the chloride is replaced by hydroxide to pro-

Haemoglobin. The pigment of the red blood cells which is responsible for the transport of the oxygen from the lungs to the tissues. It is a conjugate protein composed of the iron-porphyrin compound of haem, combined with the basic protein globin. It

combines with oxygen in the lungs. It also combines with CO. forming carboxyhaemoglobin. It also regulates the acidity (pH) of the blood, acts as a respiratory catalyst and a CO, carrier.

Haemolysis. A phenomenon involving the bursting of blood corpuscles. It may be caused by lowering the osmotic pressure of the solution surrounding the corpuscles, by repeatedly freezing and thawing of blood and by heating to 60°C.

Hafnium, Hf. At. No. 72, At. wt. 178.6, M.P. 2220°C, D 13.3. A transition metal found associated with zirconium ores and belongs to IV group. The electronic configuration is 6s25d1. It is a strong neutron absorber and is used in nuclear reactors. It forms alloys with W and Ta.

Hair Hygrometer. An apparatus for the determination of humidity.

Hahnium, Ha. A post actinide element and has 105th position in periodic table.

Halazone, C2H5Cl2. M.P. 213°C. A white powder used for the sterilization of drinking water.

Half Cell. A single electrode in contact with a solution of ions. A potential is developed between the electrode and solution. The electromotive force of a half cell is measured by coupling it with another half cell e.g., hydrogen electrode or glass electrode.

Half-life, Half Value Period (tv2). The time taken for the concentration of a substance to fall to its half value. For a radioactive element, it is the time taken for the half the nuclei to decay and

is equal to $0.69 \times \frac{1}{\lambda}$ where λ is the decay constant.

Half Reaction. Describes the stoichiometry of the reaction of one of the reactants involved e.g. $Cr_2O_7^{2-} + 14H^+ + 6e \rightarrow 2Cr^{3+} + 7H_2O$.

Half Wave Potential, E1/2. A characteristic of a substance under investigation which is represented by the midpoint of the polarographic wave of an electrochemical reversible reaction.

Halide. A general name for fluorides chlorides, bromides and iodides.

Haloalkanes (alkyl halide). A group of organic compounds in which one or more hydrogen atoms of an alkane have been replaced by halogen atoms e.g., chloroethane (C₂H₅Cl).

Haloamines. A general name for halogen derivatives of ammonia e.g., chloramine (ClNH₂), difluoroamine (F₂NH), etc. All the

haloamines are explosive.

Trihalogen derivatives of methane represented by Haloform. general formula CHX₈ where X is a halogen atom (F, Cl, Br, 1) e.g., CHF₈, CHCl₈, CHBr₈ and CHI₈.

- Haloform Reaction. A reaction used for the chemical recognition of methyl carbinol and methyl carbonyl structures which forms CHCl, with NaOCl, etc.
- Haloform Test. A test for detecting any of the halo forms or dihalocarbenes. The test is carried out by adding 10% NaOH and pyridine to haloform resulting in the formation of pink to
- Halogenation. A process of introducing halogen atoms in to molecules by addition or substitution e.g., formation of chloromethane (CH₃Cl) from methane (CH₄) and chlorine (Cl₂) or the formation of chlorobenzene (C₆H₅Cl) from benzene (C₆H₆) and chlorine (Cl₂) in the presence of aluminium chloride catalyst.

Halogen. A group of elements fluorine, chlorine, bromine, iodine

Halothane, Fluothane, CHBrCl.CF3. B.P. 45-51°C. A non-inflammable heavy liquid used as a general anaesthetic mixed

Hammer Mill. A type of mill which operates by impact.

- Hammick and Illingworth's Rule. A rule governing the course of aromatic substitution. If C₆H₅X₄ represents a monosubstituted benzene derivative where X is attached directly to the nucleus and Y is an atom or group attached to X, then if:
 - (a) Y is in a higher group in the periodic table than X₁ or
 - (b) Y is in the same group as X but of lower atomic weight, XY is meta directing. But if
 - (c) Y is in a lower group than X, or
 - (d) X and Y are atoms of the same element, or
 - (e) XY is just a single atom, XY is ortho and para directing.

Hydrogen should be considered as a member of I group: in such a case as CHO the hydrogen must be ignored and only the

- Hantzsch Synthesis. A reaction used for the preparation of pyridine derivatives (also pyrrole) by the condensation of ethylacetoacetate
- Haptens. Substances responsible for specificity of many antigens.
- Hapto. The number of atoms in a ligand which are bonded to
- Hardening (of oils). A process used for converting liquid vege-
- Hardness (of water). A property of water that will not form a lather with soap due to the presence of dissolved Ca, Fe and

A property of a material depending on the pressure applied to small area e.g., resistance to crushing.

HCH. Hexachlorocyclohexane.

hep. Hexagonal closed packing.

He. Helium.

Heat Engine. A device for converting heat energy into work.

Heating Oil. Oils used in domestic and industrial heatings e.g gas oils and kerosines.

Heat of Atomization. The amount of heat required to dissociate one I gm molecule of an element into its atoms.

Heat of Combustion. The amount of heat evolved when 1 gm molecule of the substance is burnt in oxygen at constant volume.

Heat of Crystallization. The heat evolved when unit weight of solute is crystallized from a large quantity of saturated solution.

Heat of Dissociation. The amount of heat required to dissociate 1 mole of a compound into its elements, or into certain specified smaller molecules.

Heat of Formation. The heat energy changed (evolved or absorbed) when I mole of substance is formed from its elements at I atm. pressure and the specified temperature. The standard heat of formation of an element at standard state is taken as zero.

Heat of Neutralization. The amount of heat evolved when one mole of an acid or base is completely neutralized.

Heat of Reaction. The amount of heat absorbed or evolved when specified amounts (I mole of reactant react completely at a constant pressure). The enthalpy change for exothermic reactions is taken as negative e.g., $\triangle H = -ve$, while for endothermic reactions, $\triangle H = +ve$.

The heat of reaction is generally expressed in kilojoules (kJ).

Heat of Solution. The heat evolved or absorbed when one mole of a substance is dissolved in a given solvent to form a dilute solution.

Heat Transfer Media. Fluids used to transfer heat from the place of generation to the required location.

Heat Transfer Oils. Oils used in the transformation of heat from one place to another.

Heavy Hydrogen. Deuterium; D; 2,1H.

Heavy Spar. BaSO4.

Heavy Oil. An oil obtained from the fractional distillation of coal tar.

Heavy Water (deuterium oxide), D₂O. The residual water obtained from the electrolysis of dilute aqueous solutions of acids or alkalis. Used as a reaction absorber, or "moderator", in the atomic energy processes and as tracers.

Heisenberg Uncertainty Principle. It is impossible to determine accurately both the position and momentum of small particles which possess both wave and particle properties. Mathematically.

$$\triangle p \times \triangle x = \frac{h}{2\pi}$$

where $\triangle p$ is the momentum, $\triangle x$ is the uncertainty in the position and h is Planck's constant.

Helium, He. At. No. 2, At. wt. 4.008. Discovered by Jansen. colourless monoatomic gas, the first member of noble gases (group 0 of the periodic table) with electronic configuration 1,2. Its nucleus contains 2 protons and 2 electrons with two extranuclear electrons. Found in radioactive minerals as a product of radioactive decay. Occurs in some natural gas (7%) and is separated from natural gas by liquefaction. Used in the field of inertness, e.g., as an inert atmosphere for arc welding and for Ti, Zr, Si, Ge production, as a coolant in nuclear reactors, as an atmosphere for divers, etc., liquid. He has no triple point. 2.2°K liquid. He undergoes a transition from liquid He, to liquid HeII, the later being a true liquid which exhibits conductivity.

Hell-Volhard-Zelinsky Reaction. A method for the preparation of halogenated carboxylic acids using free halogen in the presence of phosphorus halide catalyst.

RCH₂·COOH→RCHBr.COOH

Helmholtz Free Energy, F. A thermodynamic function which can be defined as the maximum amount of energy available to do work resulting from changes in a system at constant volume.

F = U - TS

where U is the internal energy, T is the thermodynamic temper-

Hemimellitene, C₆H₃(CH₃)₃. 1, 2, 3-trimethoxy benzene.

Hemimorphite, (OH)₂ Zn₄ Si₂O₇, H₂O. An ore of sinc.

Henderson-Hasselbach Equation. An equation used to calculate

where Ka is the dissociation constant of the acid component of

Henry, H. The SI unit of inductance and is equal to the inductance of a closed circuit that has a magnetic flux of one weber per

IH=IWbA-1

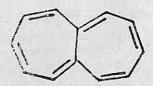
Henry's Law. The mass of a gas (m) dissolved by a unit volume of a given liquid is proportional to the pressure (p) of the gas at a constant temperature.

i.e.,
$$\frac{m}{p}$$
 = Constant = k

where p is the partial pressure of the gas in equilibrium with the solution, c is the concentration of the gas (molar) and k is a constant of proportionality. Also, the volume of the gas absorbed in unit volume of a given liquid is independent of pressure.

Heparin. A compound obtained from liver that prevents the blood clotting and is used to treat thrombosis.

Heptalene, C12H10. An anti-aromatic hydrocarbon.



n-Heptane, C7H16, CH3.(CH2)5.CH3. A colourless liquid alkane obtained from petroleum refining. Used as a solvent.

Heptose. A hydrocarbon containing 7 carbon atoms.

Herbicides. Weed killers.

Heroin, C21H23NO5, diamorphine. A morphine salt, highly addictive and used in the form of hydrochloride which is soluble in water and alcohol. M.P. 232°C.

Hertz, Hz. The SI unit of frequency, defined as one cycle per second. Applied to vitaration or wave motion.

Hess's Law (of constant heat summation). This law states that for a given chemical process, the net heat change $(q_p \text{ or } q_v)$ will be for a given chemical process occurs in one or several stages (independent of rate of the process). It is an application of first law of thermodynamics to chemical reactions.

Hetero Atom. See heterocyclic compounds.

Heterocyclic Compounds. Organic ring compounds containing more than one type of atom, e.g.,

pyridine (), pyrrole (), etc.

Heterogeneous. Describing more than one phase, e.g., water in contact with its vapours has two phases (liquid and vapour).

Heterogeneous Catalysis. A reaction in which the catalyst is in different phase than that of reactants, e.g., V2O5 catalyst in contact process for the manufacture of sulphuric acid.

Heterogeneous Reaction. A reaction taking place between different phases, e.g., a gas and liquid.

Heteroleptic. A compound containing more than one substituent,

Heterolytic Fission. The breaking of a covalent bond in such a way that both the electrons of the linkage are retained by one species $RX \rightarrow R^+ + X^-$

Heteropoly Acids. A group of complex acids (e.g., phosphotungstic acid) having anious which are derived from two different acid-

Hexaborane, B₆H₁₀ and B₆H₁₂. Boron hydrides.

Hexachlorobenzene, C₆Cl₆. Colourless crystals, M.P. 227°C. Prepared from C₆H₆ and Cl₂. Used as a wood preservative.

Hexachloroethane, C₂Cl₆. Colourless solid, M.P. 187°C.

Hexadecane Cetane, $C_{16}H_{34}.CH_{3}[CH_{2}]_{14}.CH_{3}$. M P. 18°C. An alkane.

Hexadecanoate, (palmitate). A salt or ester of hexadecanoic acid.

Hexadecanoic Acid (palmitic acid), CH₃(CH₂)₁₄COOH. A carboxylic acid present in fats and oils as glycerides.

Hexagonal Close Packing. An important way of packing equal spheres to occupy the minimum volume per sphere. The co-ordination number is 12. Zn and Mg show hexagonal close-packed structures.

Hexahelicene, C₂₈H₁₆.



Hexametaphosphate. Derivatives of Graham's salt (NaPO₃)_n.

Hexamethylbenzene, C₁₂H₁₈. Colourless crystal, M.P. 164°C.

Hexamine, C₆H₁₂N₄. A white crystalline solid obtained from methanal and ammonia. Used as a fuel, in the manufacture

Hexane, C₆H₁₄. A liquid alkane obtained from the light fraction of crude oil. Used as a solvent.

Hexanoic Acid (caproic acid), CH3(CH2)4COOH. An oily carboxvlic acid present in cow's milk and same vegetable oils.

Hexabarbitere, C,2H,6N2O3. A rapidly absorbed hypnotic. Used to produce general anaesthesia.

Hexose. A sugar containing 6 carbon atoms and a building unit of almost all polysaccharides, e.g., D-glucose, D-mannese (aldohexoses) and fructose, sorbose (ketohexoses).

Hexyl. The group CoH13.

Heyrovsky-Ilkovic Equation. An equation used in polarography.

$$E_{DME} = E_{\frac{1}{2}} + \frac{0.059}{n} \log \frac{i_d - i}{i}$$

where E1/2= half wave rotential, i=current, EDME=the potential at the mercury dropping electrode, n=no. of electrons and ia=the diffusion current.

Hf. Hafnium.

Hg. Mercury.

High Spin Rate. A term describing the transition metal chemistry that the compound has the maximum number of unpaired electrons consistent with the eletronic configuration.

High-temperature Carbonisation. A process of treating coal, in the absence of air and at a temperature from 800° - 1100°C for about 10-24 hours.

Hippuric Acid, Benzoylglycine, C9H9O3N. M.P. 187°C, soluble in water and alcohol. Excreted in the urine of mammals.

Hirudin. A substance obtained from the salivary gland.

Histamine, C. HoN3,

A base, present in animal tissues and ergot.

Histidine, α-Amino-β-imidazolpropionic Acid, C₆H₉O₂N₃. Colourless plates. M.P. 277°C, soluble in 24 parts of water. An essential constituent of the food of animals and one of the basic amino acids occurring in the hydrolysis product of proteins.

Histones. A class of proteins, found in combination with nucleic acid and are present in red blood corpuscles and in spermatozoa.

HMPA, HMPT. Hexamethylphosphoric triamide.

Hoesch Synthesis. A reaction widely applied to the synthesis of anthocyanidins by condensing polyhydric phenols with nitriles by the action of HCl.

Hofmann Degradation. A method of preparing amines from acid amides. The amine is obtained by refluxing amide with aqueous NaOH and Bra.

 $RCONH_2+NaOH+Br_2\rightarrow RCONHBr+NaBr+H_2O$ $RCONHBr + OH^{-} \rightarrow RCON^{-}Br + H_2O$ $RCON^{-}Br \rightarrow R - N = C = O + Br^{-}$ $RNCO+2OH^{-}\rightarrow RNH_2+CO_3^{2-}$

Hofmann's Method. A process for determining the vapour density

Holmium, Ho. At. No. 67, At. wt. 164.93. An element of the rare earth group which is associated with some lanthanides.

Homo. A prefix used in organic chemistry and is due to a difference

Homocyclic Compounds. Compounds containing a ring made up of the same atoms, e.g., C₆H₆ (benzene).

Homogeneous Catalysis. A chemical reaction in which the catalyst and the reactants are in the same phase, e.g., the acid catalysed

Homogeneous Reactions. Reactions in which all the reactants are in the same phase, e.g., the synthesis of ammonia from nitrogen and

 $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$

Homoleptic. Compounds having only one type of substitution.

Homologous Series. A group of organic compounds having the same functional group and a regular constitutional pattern with a difference of fixed number of atoms. The members of a homologous series can be represented by a general formula. For example, alkane can be represented by a general formula

 $C_nH_{2^{n}+2}(CH_4, C_2H_6, C_3H_8,....).$

Any two successive members of a series are known as homologues.

Homolytic Reaction (fission). A process involving the breaking of a covalent bond such that each species formed possesses one electron, resulting in the formation of free radicals.

Homonuclear Molecule. A molecule made up of identical atoms of

Homopolar Bond. A covalent bond.

Homopolar Crystal. A crystal having all covalent bonds, e.g.,

Hormones. Compounds secreted directly into the blood stream by some organs of the body (endocrine gland), e.g., thyroxine, insulin etc.

Hot Carbonate Process. A process for removing CO2 from gases by extraction with hot solutions of NaOH or K2CO3.

HPLC. High pressure liquid chromatography.

Humectant. A substance used to preserve the moisture content of materials, e.g., glycerol.

Hume-Rothery's Rule. The ratios of the total valency electron to the number of atoms in the empirical formula determine the phase of some alloys.

Humic Acid. A group of high molecular weight organic acids.

Humidity. An absolute humidity is the mass of water vapour per unit mass of dry air.

The relative humidity is the ratio of the partial pressure of the water vapour in the air to the partial pressure of the water vapour in the air when saturated at the same temperature (expressed as a percentage).

Humus. A dark coloured, amorphous organic constituent of

Hund's Rule (of maximum multiplicity). In a given subshell electrons avoid entering the same orbital and the electrons enter ing the different orbitals of a sub-shell have parallel spin, e.g., p^3 subshells will have electrons: $(\uparrow)(\uparrow)(\uparrow)$ rather than $(\uparrow\downarrow)(\uparrow)$.

Hybridization. A process in which dissimilar orbitals of nearly the same energy on the same atom may combine to from an equal same energy on the same are number of equivalent energy new orbitals. The new orbitals are said to be hybrid orbitals.

Hydantoin, Glycolylurea, C3H4N2O2

HN

M.P. 220°C. Colourless needles soluble in alcohol and present in beet molasses.

Hydnocarpic Acid, C₁₆H₂₈O₂. A fatty acid. Hydracrylic Acid, β-Lactic Acid, 2-Hydroxy Propionic Acid, CH₂OH.

CH₂COOH. A syrup, used in the manufacture of esters. Hydrates. Compounds having molecules of water of crystallization,

Hydraulic Fluids. Fluids used in hydraulic press.

- Hydrazides. The alkali metal derivatives of hydrazine, e.g., NaNH.NH2 (Sodium hydrazide).
- Hydrazine N₂H₄, (H₂N.NH₂). M.P. 14 C, B.P. 114 °C. A colourless liquid obtained by the oxidation of ammonia with NaClO₃ or by the gas phase reaction of ammonia with chlorine. It is a weak base and forms salts with strong acids (e.g., N₂H₄.HCl). Used to form various derivatives of crganic compounds, as a fuel for jet engines and for rockets.
- Hydrazoic Acid, N₃H (hydrogen azide, azoimide). M.P. -80°C. B.P. 37°C. A colourless liquid prepared from sodium azide and dilute acid (distillation). It is highly poisonous and explodes in the presence of O2 and other oxidising agents. Generally used in aqueous solutions for various synthesis in organic chemistry. Lead azide

Hydrazones. A type of organic compounds C=N.NH₂ obtained

by condensing an aldehyde or ketone with hydrazine. Phenyl hydrazine (C₆H₅NH.NH₂) forms phenyl hydrazones. 2, 4-dinitrophenyl hydrazine is used for preparing crystalline derivative for identification of keto-compounds.

Hydrides. Compounds of hydrogen. The important classes

- (a) salt-like, e.g., NaH
- (b) covalent, e.g., CH₄ or H₂Fe(CO)₄ [transition metal (c) complexes, e.g., BH4-

Hydroicdic Acid, HI. An aqueous solution of hydrogen iodide.

Hydrocarbon Resins. Thermoplastic polymers of M. wt. less than 2000 obtained as a by-product in petroleum cracking, e.g., petroleum resins, terpene resins, cyclopentadiene resins, etc.

Hydrocarbons. Compounds of carbon and hydrogen, e.g., methane

Hydrochloric Acid. HCl. A colourless fuming liquid obtained from $HCl(g)+H_2O(I)-\rightarrow H_3O^++Cl^-$

It shows the typical properties of a strong acid. Used in the manufacture of dyes, drugs and photographic materials. A strongest acid of hydrohalic acids series which is oxidized to Cl₂ by oxidising agents.

Hydrochlerinated Rubber. Materials used in the films.

Hydrocracking. A process in which both the catalytic cracking and hydrogenation of petroleum fractions take place (gasoline

- Hydrofining. A process of removing alkenes from light petroleum fractions.
- Hydrofluoric Acid, HF. A colourless liquid formed by dissolving hydrogen fluoride in water.

A weak acid which is used to etch glass (dissolves most silicates). Handled by monel, tellon or equipment.

- Hydrogasification. A process converting crude oil or oil fractions to substitute natural gas (SNG).
- Hydrogen, H. At. No. 1, At. wt. 1 00794, M.P. —259 14°C, B.P. —252 87°C. A colourless lighest gas, occurs abundantly in universe and in the form of water. It is placed in zero group of the periodic table and shows similarities to both the alkali metals (group I) and the halogens (group VII). Prepared commercially by the electrolysis of water. Occurs as H₂; explodes with O₂ on sparking and reacts with halogens. Forms metal hydrides. Used in the manufacture of ammonia, for hardening edible oils, etc., Forms isotopes, e.g., deuterium.
- Hydrogenation. A process in which hydrogen is added to a substance in its gaseous form and in the presence of a catalyst. The formation of ammonia by Haber's process or hydrogenation of unsaturated liquid vegetable oils are the examples of hydrogenation.
- Hydrogen Bond. An intermolecular bond which involves bridging through a proton or hydrogen attached to an electronegative element. The length of a hydrogen bond is specifically 0.15-0.24 mn. Compounds having hydrogen bonding show high boiling points, e.g., H₂O, HF.
- Hydrogen Bromide, HBr. M.P. –88°C, B.P. –67°C. A colourless gas which is quite soluble in water. Obtained by the direct combination of H₂ and Br₂ in the presence of a platinum catalyst. Dissolves in water to form hydrobromic acid. A molecular compound which conducts electricity in molten state.
- Hydrogen Carbonate (bicarbonate. A salt containing the ion
- Hydrogen Chloride, HCl. A colourless fuming gas, obtained by the action of concentrated sulphuric acid on NaCl. Forms dense white fumes with ammonia. Highly soluble in water and ionizes white fumes with accordance acid. It has a strong H-Cl bond and to produce hydrochloric acid. It has a strong H-Cl bond and is used in the manufacture of organic chlorine compound, e.g., polyvinyl chloride (PVC).
- Hydrogen Cyanide, Hydrocyanic Acid, Prussic Acid, HCN. M.P. -13°C, B.P. 26°C. Colourless, miscible with water, burns in air, smell of bitter almonds. Obtained from CH₄, air, NH₃ in the presence of a catalyst at 1000°C. Forms salts, cyanides, isoni-

triles, etc. Polymerises on standing. Used as a funmigant and

Hydrogen Electrode. A reference electrode (standard electrode) used to assign numerical values of potentials to other electrodes. The electrode is the normal hydrogen electrode (NHE). H₂(Pt) gas

H⁺(a=1) in which pure hydrogen gas at unit pressure is kept in contact with solution containing H+-ion of unit activity through adsorption on Pt-black by continuous bubbling of the gas. The potential of this hydrogen electrode is taken as zero at all temperatures. To evaluate the potential of any other single electrode, it is coupled with NHE and the emf of the galvanic cell is measured potentiometrically. Mathematically,

E_{H₂}=E_{H⁰₂}-
$$\frac{RT}{nF}$$
 ln $a_{H+}=-\frac{RT}{nF}$ in a_{H+}
if $a_{H+}\neq 1$; $E_{H_2}\neq 0$.

where $E^0_{H_2}$ =the oxidation potential of hydrogen electrode. If the given electrode functions as negative electrode, then

 $E = E_{ano.le} - E_{cathode} = E_x - 0 \quad [since \quad E^{0}_{H_2} = 0]$ given electrode functions as positive electrode, then and, if the

 $E = E_{anede} - E_{cathode} = E^{0}_{H_2} - E_x = -E_x$

where Ex is the oxidation potential of the other electrode (given

Hydrogen Fluoride, HF. M.P. -83°C, B.P. 19.5°C. A colourless, strongly fuming, associated liquid. Prepared from KHF2 or CaF₂ and H₂SO₄. CaF₂(s)+H₂SO₄(aq)→CaSO₄(aq)+2HF(l)
Association of H-F units is due to the presence of hydrogen bonding in hydrogen and highly electronegative fluoride ions. Used in the preparation of Al, fluoro carbons, steel manufacture and as a catalyst in petroleum industry.

Hydrogen Halides. HF, HCl, HBr and HI, only HF is associated molecule containing strong hydrogen bonding.

Hydrogen Iodide, HI. M.P. -54°C, B.P. -36°C. Colourless gas, soluble in water. Prepared from H₂ and I₂ in the presence of a

Hydrogen Ion Concentration. It is the number of gram ions present per litre of solution. As these concentrations are very small, the hydrogen ion concentration is generally expressed as the pH of the solution which is the negative logarithm of hydrogen

 $pH = -\log C_{H^+}$ or $pH = -\log a_{H^+}$ where a_H^+ =activity of hydrogen ions.

Hydrogen Over-voltage. In electronic processes Caspari observed the potential at which the visible evolution of H₂ and O₂ gases occurred at cathodes and anodes, respectively, of a number of different metals. The difference between the potential of the electrode when gas evolution (H₂) was actually observed and the theoretical reversible value for the same solution was called hydrogen over-voltage.

Hydrogen Peroxide, H_2O_2 . M.P. -0.4° C, B.P. 150°C, D 145. A colourless syrupy liquid, with an odour like that of nitric acid. Evaporates spontaneously in air and forms a crystalline hydrate, $H_2O_2.2H_2O$. Prepared by oxidation of anthraquinone with O_2 followed by reduction of quinone with hydrogen. Decomposes in aqueous and alkaline solutions to oxygen and water (catalysed by dust). Pure H_2O_2 in stable but in contact with manganese (IV) oxide decomposes to form O_2 and water.

 $2H_2O_2 \rightarrow 2H_2O + O_2$

Often used in aqueous solutions. Concentration expressed in volume as volume strength—the volume of oxygen (dm³) at STP given by decomposition of 1 dm³ of the solution. H₂O₂ is generally used in 3% (10 volumes), 6% (20 volumes), or 30% solution (100 volumes). A 3% solution forms 10 times its volume of oxygen on dissociation. A powerful oxidizing agent. Used in chemicals, textiles, paper and as a propellant. Also used as a medicine.

Hydrogen Persulphides, Sulphanes, H_2S_x (x=2-6). Obtained from polysulphides and acids followed by cracking. Decomposes to H_2S and S.

Hydrogen Selenide, H₂Se, M.P. -64°C, B.P. -42°C. A colourless gas prepared by the action of aluminium selenide on 5N-HCl.

Hydrogen Sulphide, Sulphuretted Hydrogens, H₂S. A colourless gas with unpleasant odour of rotten eggs and very poisonous. Occurs in some mineral waters and also produced by decaying animal or vegetable matter. Prepared from FeS and dilute H₂SO₄. Aqueons solution is weakly acidic.

 $H_2S \rightleftharpoons HS^- + H^+$ $HS^- \rightleftharpoons H^+ + S^2^-$

Forms insoluble sulphides with most metal sulphides in solution (used in qualitative analysis of salt mixtures). Burns with a blue flame in oxygen to produce sulphur (IV) oxide. Detected by lead acetate which forms black precipitate with H₂S.

Hydrogen Telluride, H₂Te. A colourless solid, M.P. -57°C, B.P. 1'8°C. Soluble in water and ether.

Hydrolith, CaH2. Calcium hydride.

Hydrolysis A process of interaction between chemical compounds and water. When salts are dissolved in water the resulting solution is generally, either acidic (if free H⁺ ions are present in solution), in solution) or basic (if free OH⁻ ions are present in solution),

e.g., ammonium chloride solution is acidic while sodium carbonate solution is basic in nature.

$$NH_4^+ + H_2O \rightleftharpoons (NH_4^+ + OH^-) + H^+$$
weak base

$$CO_3^{2-} + H_2O \rightleftharpoons HCO_3^- + OH^-$$

The hydrolysis constant (K_h) is given by:

$$K_h = \frac{[NH_4OH][H^+]}{[NH_4^+][H_2O]}$$

Since, water is present in excess, so (H2O) is taken as constant,

$$K_h = \frac{[NH_4OH][H^+]}{[NH_4^+]}$$

In organic chemistry the term is applied to the hydrolysis of

Hydrometer. An apparatus for measuring the density or specific

Hydroperoxide. A group of compounds containing the hydroperoxyl group, HO.O-, e.g., H₂O₂.

Hydrophilic. Water attracting.

Hydrophilic Colloid. Colloidal sols of proteins, carbohydrates, soaps, etc., which are not coagulated by small amounts of electrolytes such as are required to coagulate hydrophobic or suspensoid sols (e.g., gold sol). The stability of such sols is due to the presence of a sheath of dispersion medium (water)

Hydrophobic. Water repelling.

Hydrophobic Sol. Sols of gold, arsenic sulphide, etc., which are stable due to the mutual repulsion of like charges on all the colloidal particles. They are not hydrated and coagulated by a small amount of electrolyte (due to neutralisation of charge on sol). They are coagulated only at the iso electric point.

Hydroquinone, C₆H₆O₂, colourless solid, M.P. 170°C. Used as a

Hydrosulphurous Acid, H2S2O4. Dithionic acid.

Hydroxide. A corepound having OH- ion or the group OH.

Hydroxonium Ion, [H (H2O)n]+.

Hydroxyapatite, Ca5 (OH) (PO4)3.

Hydroxylamine, H2N.OH. M.P. 33°C. A colourless solid, forms stable salts; e.g., hydroxylamine hydrochloride (NH2OH.HCI). Acts as a reducing and oxidizing agent. Used in the production of nylon,

Hydroxylation. The process of introduction of hydroxyl group

into an organic compound.

Hydroxyl Ion. The OH ion.

2-Hydroxypropionic Acid, Lactic Acid, CH3CH(OH)COOH. A colourless liquid.

Hydroxy Propanone, Acetol, Hydroxy Acetone, Pyruvic Alcohol, CH₈CO. CH₂OH. B.P. 145-146°C. Colourless liquid which tends to polymerize.

8-Hydroxy Quinoline, Oxine, CoH7ON. M.P. 75-76°C. Light brown solid. Forms Cu derivates which are used as fungicides.

Hygroscopic. Describing a substance which absorbs moisture from atmosphere, e.g., MgCl₂ or CaCl₂. Used as desiccants.

Hyoscyamine, C17H23NO4. Highly poisonous syrup. Used as a sedative in small quantities.

Hyperconjugation. The magnitude of the effect in which alkyl groups interact electronically with unsaturated substances to which they are attached, in a similar way to unsaturated groups.

Hyperons. Elementary particles present in cosmic radiation. Mass of a hyperon is 2185 times that of electron.

Hypo, Na₂S₂O₃. Sodium thiosulphate. Used in photography.

Hypobromous Acid, HOBr.

Hypochlorites, M[ClO]. Salts of hypochlorous acid.

Hypochlorous Acid, HOCl. Used as bleaching agent and powerful

Hypofluorites. Covalent compounds having -OF groups, e.g., F2Q, SF5OF, etc.

Hypophosphoric Acid, H₄P₂O₆. A tetrabasic acid obtained from NaClOo and red phosphorous.

Hypophosphorous Acid, H₄P₂O₆. M.P. 26°C. A monobasic acid pophosphorous Acid, H_4 206. heating with NaOH. Acid and which liberates H_2 and PH_3 on heating with NaOH. Acid and its salts (e.g., NaH₂PO₂) are powerful reducing agents.

Hyposulphurous Acid, H2S2O4. Dithionous acid. Hysteresis. A term applied to solids, e.g., during the process of magnetization of iron.

in plants and in human body.

yellow solid, soluble in water and alcohol.

Inductive Effect. The effect in which atoms or groups in a compound can attack (-I) or push away electron (+I) forming polar bonds. For example, trichloroethanoic acid (CCl₃.COOH) is much acidic than ethanoic acid CH₃COOH.

Industrial Methylated Spirit, IMS. Ethanol containing wood

Inelastic Neutron Scattering. Describing a process involving the bombardment of a crystal by a beam of monochromatic neutrons resulting in the scattering of the beam due to collision with crystal nuclei. The process provides information about the crystal structure and vibration within the crystal.

Inert Atmosphere Box. A box used for chemical reaction in the absence of water, oxygen and nitrogen.

Inert Complex. A complex that takes part in reactions very slowly so that the rate of reaction can be easily studied. Inert Gases. Noble gases.

Inert Pair Effect. A co-relation between the oxidation states n and (n-2) and the s^2 co-relation for the latter for the heavy p block

Infrared, (IR). Electromagnetic radiation with longer wavelengths than visible radiation (frequency less than 1013 per second and wavelength greater than about 8000A°). Invisible to the human eye with adequate penetrating power. IR radiations are applied in long distance photography, in night viewing relay on detection

Infra-red Spectroscopy, (IR spectroscopy). IR radiation is produced by vibrational or rotational motion of molecules (within). These motions show characteristic absorption bands in the IR region of spectrum. Certain bonds between pairs of atomsquencies, which correspond to band in IR spectrum. IR spectroscopy provides valuable information about molecular structure, groups present and molecular symmetry. They are also used to identyfy new compounds (fingerprint).

- Infusions. Dilute solutions of some drugs.
- Ingrain Dyestuffs. Insoluble dyestuffs formed directly on the fibre from soluble components.
- Inhibitor. An additive which will inhibit, i.e., slow down or stop a reaction. For example, H₂S, HCN, Hg-salts and As compounds inhibit heterogeneous catalysts by adsorption. An inhibitor acts by preventing the propogation of chain (e.g., chain reactions) or by passivating the surfaces (e.g., corrosion reactions).
- Ink. Colouring matter which has been dissolved or dispersed in a solvent and is bonded to the surface on drying (a colloidal sol).
- Inorganic Chemistry. The branch of chemistry dealing with all the elements other than carbon. Certain simple carbon compounds constitute a part of inorganic chemistry. For example, carbide, carbonates, carbonyl, cyanides, etc.

Isosinic Acid, C₁₀H₁₃N₄O₈P. A purine nucleotide.

Inositol, C6H12O6. Hexahydrocyclohexane.

Insecticides. Material used to control insects, e.g., CS2.

Instability Constant. A constant and measure of dissociation.

Insulators. Substances with a very poor electrical conductivity, e.g., pure solid ionic substances.

- Instrumentation. The measurement of the conditions and the control of processes with in a chemical plant.
- Insulin. It is built up of two polypeptide chains, 'A' of 21 amino acids and 'B' of 30 amino acids, linked by two disulphide acids and 'B' of secreted by the islets of Langerhans in the bridges. A hormone, secreted by the islets of Langerhans in the pancreas. The absence of insulin leads to the disease diabetes.

Interatomic Distances. The distance between the nuclei of atoms (bonded or non-bonded) in a molecule or crystal. Determined by spectroscopy, electron diffraction method, etc.

Intercalation Compounds. A compound particularly of graphite (a layer structure) in which molecules are accommodated between the layers in lattice or in holes. For example, muscovite (KAI₂ the layers in lattice or in holes. For example, muscovite (KAI₂ the layers in lattice or in holes. For example, muscovite (KAI₂ the layers in lattice or in holes. For example, muscovite (KAI₂ the layers in talc and pyrophyllite respectively or graphite—silicon layers in talc and pyrophyllite respectively or graphite—SbF₅.

Interfacial Angles, Constancy of. The angle between the adjacent and is independent of the size of the crystal.

Interferon. A protein with lower molecular weight.

- Interhalogen Compounds. Compounds formed by the halogen elements with one another. Four important classes of such compounds are :
 - (a) AB, e.g., CIF, BrF, etc.
 - (b) AB3, e.g., CIF3, BrF3, ICl3
 - (c) AB5, e.g., BrF5 and IF5
 - (d) AB7, e.g., IF7,

The anion [IBrCl] is a polyhalide ion.

- Internal Compensation. Optical activity in which the two asymmetric atoms of the compound produce equal and opposite rotation of the plane polarized light resulting in zero optical rotation.
- Internal Energy, E or U. A thermodynamic term related to the

H=E+PV

where P is the pressure and V is the volume. Also defined as the total energy of a system, i.e., the sum of kinetic and potential energies of the constituent particles of a system.

- Interstitial Compounds. Compounds in which an atom moves off its lattice point to an interstitial position. An interstitial is an atom that is in a position that is not normal lattice point. For example, introduction of H,B,C,N, etc., into the holes of a metallic lattice (usually a transition metal).
- Intrinsic Energy. As arbitrary assumption used in thermochemistry that the intrinsic energies of all the elements is zero. intrinsic energy of a compound is equal to its heat of formation
- Invar. An iron alloy containing 35-36% Ni, 0.5% C, and 0.5% Mn which has zero co-efficient of thermal expansion up to
- Invariant System. A term often applied in phase diagrams. system in equilibrium is said to invariant when it has no degree
- Inversion. A change from one optical isomer to the other.
- Inversion Temperature. The majority of gases are generally cooled by rapid expansion (The Joule-Thompson effect). Below -80°C hydrogen is cooled but above this temperature it is warmed by rapid expansion. This temperature is known as inversion temperature.
- Invertase, Sucrose. An enzyme that hydrolyses sucrose to glucose

Invert Sugar. A mixture of glucose and fructose (1:1) obtained by hydrolysing sugar. Sucrose is dextrorotatory, $[\alpha]b = +66.5$ whereas invert sugar is laevorotatory, $[\alpha]b = -20.6$.

Todates. Salts of iodic acid (HIO3) obtained by dissolving iodine in aqueous alkalis. For example,

 $3I_2+6KOH\rightarrow 5KI+KIO_3+3H_2O$.

They are strong oxidizing agents (potential)

 $[IO] \rightarrow I + 0.49 \text{ volts}$;

 $[IO_3] \rightarrow I + 1.085 \text{ volts}$;

and $H_5IO_6 \rightarrow [IO_3]^- + 1.7$ volts.

- Iodic (V) Acid, HIO₃. A deliquescent crystalline solid and a strong oxidizing agent. Obtained from concentrated HNO₃ and iodine. Liberates iodine from iodide ions solution and produces flames with organic materials. A strong acid.
- lodic (VII) Acid, Periodic Acid, H5IO6. White solid obtained by low temperature electrolysis of HIO_3 . A powerful oxidizing agent but a weak acid. Forms dimesolodic acid $(H_4I_2O_9)$ and metalodic acid (HIO_4) . Used for the estimation of manganese in steel.
- lodimetric Method. Describing a volumetric method in which iodine is directly titrated with reagents such as, sodium thiosulphate (Na₂S₂O₃) solution.

 \rightarrow 2NaI+Na₂S₄O₆ $I_2 + 2Na_2S_2O_3$

Iodine, J. At. No. 53, At, wt. 126.92. M.P. 113.5°C, B.P. 184.4°C. Occurs in small amounts in sea water and in sea weeds. A dark violet volatile solid element of the halogen group (VII group of periodic table). Obtained commercially from the mother liquor from the extraction of sodium nitrate from "caliche'. Iodine is obtained by treating mother liquor con-taining sodium iodate) with dilute sulphuric acid and sodium hydrogen sulphide. Forms violet solutions with CCl4 or uydrogen sulphide. Forms The element is a black solid giving CHCl₃ (a test for iodine). Used as germicide in a violet vapour (I₂ molecules). Used as germicide, in medicines (for the treatment of hyperthyroidism—a disease due to disorder of thyroid gland), in photography, etc. Iodine is an

Iodine Monochloride, ICl. B.P. 97°C. A dark red liquid. Used as a

Iodine Fluorides, IF₃, Iodine Trifluoride. Yellow solid. IF₅—colourless liquid, M.P. 9°C, B.P. 100°C, used as a solvent. IF₇—B.P.

Iodine Oxides, I2O5, Iodine Pentaoxide. A white crystalline solid obtained from lodic acid (HlO₃) at 200°C. A strong oxidizing agent. Used in the volumetric titrations.

- Iodobenzene Iodobenzene, C₆H₆I. B.P. 88°C. Colourless liquid. Prepared by the iodination of C₆H₆ with iodine and nitric acid under
- Iodoform, CHI₃ (trilodomethane). M.P. 119°C. A yellow crystal-line solid with a characteristic odour. Prepared by warming ethanal with an alkaline solution of an iodide:

$$CH_3CHO + 31^- + 4OH^- \rightarrow CHI_3 + HCOO^- + 3H_2O$$
.

This reaction is given by all ketones of general formul 1 CH3COR and by secondary alcohols CH₃CH (OH) R. Manufactured by the electrolysis of a solution of an iodide in dilute alcohol or propanone. Decomposes slowly by light and air to CO₂, CO, I₂ and H₂O. Soluble in organic solvents, e.g., alcohol, ether

- Iodoform Test. A test for the detection of a CO. CH₃ group
- Iodomethane, CH₃I (methyl iodide). A liquid obtained by the action of methanol and iodine in the presence of red phos-
- Iodiometry. A volumetric method for the estimation of oxidizing agents (e.g., copper sulphate, potassium dichromate) which liberate I2 from acidified KI solutions (an indirect method).
- Ion. An atom or group of atoms which has gained or lost one or more electrons resulting in a negative or positive charge, e.g.,
- Ion Exchange. A process exhibited by insoluble materials, which contain ions capable of exchanging ions with ions in the surrounding medium, e.g., zeolites used for softening of water or sodium in certain silicates can be replaced by potassium by exhaustively washing the material with a solution of KCl. An anionic resin exchanges negative ion, when all available ions have been exchanged (e.g., Na⁺ replacing Ca²⁺ ions), the material can be regenerated by passing concentrated solution (e.g., NaCl solution) through it. Using a 'mixed bed' of anion and cations exchange resins the electrolytes can be removed from a water solution, and very pure water obtained.
- Ion Exchange Chromatography. See chromatography.
- Ion Exclusion. A process used for a partial separation of electrolytes from non-electrolytes. The medium inside the pores of an ion-exchange resin contains a lower concentration than that of the external solution of any electrolyte that may be present in the latter (Donnan membrane equilibrium). This is known as

Ionic Atmosphere. An important effect used to study the electrical conductivities of solution. According to Debye-Huckel theory, in solutions, ions carrying one type of charge are surrounded by an atmosphere of ions carrying opposite charge. Under the influence of an applied electric field, the central ion moves in one direction whereas the ionic atmosphere moves in opposite direction resulting in an asymmetric distribution of ions about the central ion.

Ionic Mobility. The velocity with which an ion would move under a potential gradient of 1 volt per centimeter in a solution is called its ionic mobility or absolute velocity. The absolute ionic velocity in cm/sec is obtained by dividing the ionic conductance by 96,500.

The ionic mobility of a cation = $\frac{1c^{\circ}}{k} = \frac{1c_0}{J} = \frac{1c_0}{96,500}$ and the

ionic mobility of an anion = $\frac{\text{la}^{\circ}}{\text{k}} = \frac{\text{la}^{\circ}}{\text{J}} = \frac{\text{lac}}{96.500}$ where

le° and la° are the equivalent conductance of cation at infinite dilution and anion at infinite dilute, respectively.

Ionic Radii. Describing the effective radius of ions in crystal lattice. Not applied to an isolated ion, since the ion is a nucleus surrounded by an 'electron cloud'.

Ionic Strength. For any electrolytic solution the ionic strength is given by

[=] Σimizi2

where m= the molal concentration and z= the charge on the ionic species.

Ionization Chamber. An apparatus for measuring the absolute intensity of a beam of X-rays or ionizing particles.

Ionization Energies, Ionization Potentials (I). The energy required to remove an electron from a free atom or ion in gaseous state. It is measured in electron volt or kilo-joules per mole. Can be represented as:

 $M \rightarrow M^+ + e$

The second ionization potential is the energy required to remove one more electron from the ionized atom.

 $M^+ \rightarrow M^{2+} + e$

The second ionization potential is always more than the first ionization potential (1eV=0.16×10-18 J).

- Ionization, Heat of. The amount of heat required to split one mole of an electrolyte into its constituent ions, e.g., the heat of ioniza-
- Ionization of Water. Pure water ionizes very slightly and exhibits a poor conductivity. In pure water,

$$[H^+]=[OH^-]=10^{-7}$$
 g ions l^{-1}

and ionic product of water, $K_w = [H^+][OH^-] = 10^{-14}$ at 25°C, but

pHof pure water is always 7.0. The pH of a solution containing H⁺ ions more than 10⁻⁷ g. ion/litre is always less than 7.0 (acidic) and a solution containing more than 10⁻⁷ g ion/litre of OH ions is always greater than 7.0 (alkaline).

- Ionogenic Complex. A complex formed on the surface of a colloidal particle which is responsible for its stability.
- Ionone, C₁₃H₂O. A powerful odorant, odour of cedar oil and forms a semi-carbazone of M.P. 148°C.
- Ion-pair. The formation of ion-pairs due to the mutual attraction of ions (contrary to Debye-Huckel theory).
- Ion selective Electrode. An electrode which responds selectively to one type of ions present in a solution and develops a potential according to the concentration of that type of ion, e.g., glass-
- Ipecacuanha. Present in the dried root of plant caphoelis ipecacuanha. Used in medicines as an emetic, in a tincture or as an
- Iridates. Irridium oxides.
- Iridium, M.P. 2410°C, B.P. 4130°C, At. No. 77, At. wt. 193'1, D 22'4. A white transition metal that is highly resistant to electronic configuration is 5d⁷ 6s² and has oxidation states from the configuration of the c $+6 (IrF_6)$ to $-1[Ir(CO_4)_4]^-$. Ir(0) generally exists as its carbonyls $Ir_2(CO)_8$. Used as an alloying material for Pt, Os and as a
- Iridium Halides. Halogen compounds of Ir, e.g., IrBr3, IrF6, IrF6,
- Iron, Fe. At. No. 26. At. wt. 55.847, M.P. 1535°C, B.P. 2750°C, D 7.87. A transition metal occurring naturally in haematite (Fe₂O₃). magnetite (Fe₈O₄). Iron pyrites (FeS₂) and chalcopyrites (CuFeS₂).

Iron is generally prepared from oxide or carbonate ores. It is extracted in a blast furnace using coke, limestone and hot air. The coke and air produce CO, which in turn reduces iron ore to iron. The impurities are removed as a slag by limestone (at the base of the furnace).

The stablest oxidation state is $+3(Fe^{3+})$ and the other state +2 (Fe²⁺) is not very stable. Ferric (+3) compounds are generally yellow whereas Ferrous (+2) compounds are green in colour.

- Iron Alums, MFe III, $(SO_4)_2$ 12 H₂O. Double salts of ferric sulphate M is an alkali metal, ammonium or univalent thallium, etc.
- Iron Ammonium Sulphate, (NH₄)₂SO₄.Fe(SO₄),6H₂O.
- Mohr's Salt. 'A light green crystalline solid used as a standard reducing agent for potassium permanganate and other oxidizing agents used in volumetric work.
- Iron Bromide, (a) Ferric Bromide, FeBr₃,6H₂O. M.P. 27°C. Dark red crystals.
 - (b) Ferrous Bromide, FeBr₂. Soluble in water, forms hydrates with 9, 6, 4 and 2H₂O.
- Iron Buff. Hydrated ferric oxide used as a black dye for silk and khaki dye for cotton.
- Iron Carbonate, FeCO₃. Occurs in nature as siderite, spathic iron ore, chalybite.
- **Iron Carbonyls.** Iron forms three carbonyls, Fe(CO)₅, Fe₂(CO)₉, and [Fe(CO)₄]₃. All a e non-polar, soluble in organic solvents and dissociate on heating to produce CO and the metal.
- Iron (II) Chloride, Ferrous Chloride. FeCl₂. A white solid obtained from Fe and HCl gas. Forms complexes, e.g., FeCl₂.6NH₃, etc.
- Iron (III) Chloride, Ferric Chloride, FeCl₃. Dark red crystals obtained from Fe and Cl₂. Forms complexes e.g., K₂FeCl₅. H₂O, (NH₄)₂ FeCl₅, H₂O, etc.
- Iron, Complex Cyanides. Derivatives of cyanoferrates e.g., nitroprussides [Fe(CN)₅NO]².
- Iron Fluorides (a) FeF₃. White salt (HF+Fe₂O₃):
 - (b) FeF₂. Obtained from Fe and HF gas. Forms hydrate and complexes such as KFeF₃.

Iron Hydroxides, Ferric Hydroxide. Naturally-occurring hydrated ferric oxides with the composition:

Fe₂O₃, ½H₂O to Fe₂O₃,4H₂O

The precipitated hydroxide is a voluminous brownish precipitate. When freshly precipitated, it is readily soluble in dilute acid. On prolonged boiling it reaches the composition Fe₂O₃, H₂O and

Colloidal, Fe(OH)3 is readily obtained as a deep red sol by partial hydrolysis of ferric chloride solution followed by dialysis.

Ferrous Hydroxide, Fe(OH)₂ is thrown down by caustic alkalis from solutions of ferrous salts (in the absence of O2).

Iron oxides produce ferrates with other metal oxides.

Iron Iodide, FeI2. Soluble in water, oxidised to I2 and ferric derivatives in air forms 9, 6, 5, 4 and 1 H₂O.

Iron Nitrate, Fe(NO₃)₃, 9H₂O and Fe(NO)₂.6H₂O.

Iron-organic Derivatives. Organic complex obtained from iron carbonyls and olefins (π-bonded complexes).

Iron Oxides (a) Fe₂O₃, Ferric Oxide. A rusty-brown solid prepared by the action of heat on Iron (III) hydroxide or FeSO₄. Occurs in nature as haematite (with corundum structure) and \(\gamma \)-form (with a structure related to spinal structure and closely related to Fe₃O₄) magnetite. Fe₂O₃ is soluble in acids with difficulty and

(b) Ferrous Oxide, FeO. A black powder prepared by the careful reduction of ferric oxide using CO or H₂. On exposure to air, it gets oxidised to Fe₂O₃. A basic oxide that dissolves

Iron Perchlorate, Fe(ClO₄)₃, 6H₂O.

Iron Phosphate, FePO₄. 2H₂O. A dark brown powder containing Iron (II) and Iron (III) phosphates and ferric hydroxides is used

Iron Sulphates (a) Ferrous Sulphate, FeSO₄.7H₂O (green vitriol). Occurs in nature as melanterite. Obtained commercially from iron pyrites and in the laboratory from iron and sulphuric acid. A freshly prepared solution of iron sulphate (ferric sulphate) absorbs nitrogen oxide (brown ring test for nitrates).

(b) Ferric Sulphate, Fe2(SO4)3. Obtained from iron sulphate (in sulphuric acid) and an oxidizing agent such as H₂O₂ or conc. HNO₃. The salt is obtained as a white mass of mono-hydrate Fe₂(SO₄)₃.9H₂O. Anhydrous salt can be obtained by gentle heating of hydrous salt. Ferrous sulphate crystals are isomeric with the sulphates of zinc, magnesium, nickel and cobalt.

Iron Sulphides. FeS2 Occurs in nature as pyrites and marcasite.

Irving-Williams Order. A general order for ligands in the following sequence

 $Mn^{2+} < Fe^{2+} < Co^{2+} < Ni^{2+} < Cu^{2+} > Zn^{2+}$.

Isatin, C₈H₅NO₂. A yellowish red powder (M.P. 200°C) used in vat dyes.

Isoamyl. The group (CH₃)₂ CH.CH₂.CH₂—

Isobars. Elements having different atomic numbers (proton+neutron) but same atomic mass (different chemical properties, i.e., different protons or electrons, e.g.,

64₂₈Ni, 64₃₀Zn and 110₄₈Cd, 110₄₆Pd; etc.

(Also lines joining points of equal pressure.)

Isobutane, 2-methylpropane, C₄H₁₀. Colourless gas, M.P. -145°C, B.P. -10°C. A by-product of petroleum cracking. Used in refrigeration.

Isobutyryl. A group (CH₃)₂CHCO.

Isocyanates. Compounds having the group—N=C=O. Obtained from dialkyl sulphates and potassium cyanates. The polycondensation products of di-isocyanates with poly-hydric alcohol are used as artificial rubber and light weight foams. Isocyanates are also used as modifiers in alkyl resins.

Isodurene, 1, 2, 3, 5-Tetramethylbenzene, C₆H₂(CH₃)₄.

Isodispersion. Describing a sol containing the dispersed particles of colloidal dimensions and all of the same size. For example, helix haemocyanate forms an isodispersion of M. wt. 5,000,000.

Isodrin. An isomer of aldrin.

Isoelectric Point. The pH at which a hydrophilic colloid has zero charge and maximum instability. At the isoelectric point the electrophoretic velocity is zero and the substance tends to flocculate.

Isoelectronic. Compounds (molecules or ions) having the same number of electrons, e.g., CO and N₂ are isoelectronic.

Isoleptic. Compounds having all ligands the same, e.g., Fe(CO)₅, [Cu(NH₃)₆]Cl₃.

Isomerase. Enzymes used to catalyse isomerization of compounds.

Isomerism. Compounds having same molecular formula but different structural formula or spatial arrangement of atoms or differing in at least one of their physical or chemical properties, are said to be isomeric and each is an isomer of the others.

- (a) Structural Isomerism—A type of structural isomerism in which the structurale formula of the compound differ. The two types of isomerism are:
- (i) Chain isomerism or nuclear isomerism—due to difference in arrangement of atoms in the molecule, e.g., butane and 2-methyl propane.

CH₃ CH₂ CH₂ CH₃ H₃C-CH(CH₃).CH₃

(ii) Position Isomerism—due to difference in position of some group or atom, e.g., C₃H₇OH represents propan-1-ol and propan-2-ol.

CH₃.CH₂.CH₂.OH and CH₃CH(OH).CH₃ propan-2-O1

(iii) Functional group isomerism or metamerism—due to the difference of a functional group, e.g., C₂H₅OH and CH₃OCH₃ or CH₃COCH₃ and CH₃CH₂CHO.

An important type of structural isomerism is keto-enol type which is exibhited by aceto-acetic ester.

 $\begin{array}{c|c}
H \\
> C - C = 0 \Longrightarrow > C = C - OH \\
\downarrow \\
(keto) & (enol)
\end{array}$

- (b) Space Isomerism or Stereo Isomerism—due to a difference in the spatial orientation of atoms in a molecule. The two important classes are:
- (i) Optical Isomerism—due to asymmetry. These isomers rotate this plane polarised light in opposite directions. Such molecules have an asymmetric atom i.e., are attached to four different groups called a chiral centre, e.g., D-lactic acid and L-lactic acid.
- (ii) Geometrical or Cis-trans Isomerism—due to a restricted rotation about a bond between two atoms, e.g., fumaric and maleic acids show cis-trans isomerism.

H-C-COOH

HOOC-C-H

fumaric acid (trans)

H-C-COOH

H-C-COOH

maleic acid (cir)

Groups attached to the same side of an atom form cis and attached to the opposite sides form trans isomer.

Some inorganic compounds also show cis-trans isomerism, e.g., [Pt(P Ph₃)_oCl_o].

Ionization Isomerism e.g.. $[Pt(NH_3)_4 \cdot Cl_2)]Br_2$ and $[Pt\ (NH_3)_4\ Br_2]$

Linkage Isomerism, e.g., the isomerism in nitrite ion.

The atomic nuclei with the same atomic and Nuclear Isomerism. mass number but different nuclear energies and radioactive decay properties, e.g., 124Sb has three isomers.

The conversion of a compound into its another Isomerization. isomer.

Isomorphism. A process in which different compounds show same crystalline state. The substances are called isomorphous.

Isonitriles, Isocyanides, Carbylamines. An organic compound having the formula R-NC. Colourless, toxic liquids, prepared by dehydration of farmamides (R NH.CHO) with phosphorus oxychlorides in the presence of a base.

R NH.CHO-→R-NC+H₂O.

A p-amine, CHCl₃ and alc. KOH forms carbylamine (a test for amines). They form stable complexes with transition metals.

Isoparaffins. Aliphatic hydrocarbons with a branched chain of carbon, e.g., isopentane.

Isopiestic. Solutions, which have the same partial pressures of solvent.

Isopolyacids. Acids which form polymeric oxide and hydroxide bridged anions having one metal only.

Isoprene, 2-Methylbutadiene, C₅H₈, CH₂=C-CH=CH₂. CH.

D²⁰ 0.6806, B.P. 34°C. A colourless liquid, insoluble in water but soluble in benzene. Manufactured from butanone and methanol in the presence of KOH to form 2-methyl-3-ketobutanol followed by reduction to 2-methylbutylene glycol. These glycol vapours when passed over heated acid phosphate form isoprene. It is converted to rubber-like substance by heat or by the action of metallic sodium, etc. Used for the preparation of synthetic rubber-like material.

Isoprene Rule. The carbon skeletons of terpenes are made up of isoprene units so that the C_4 of one unit is attached to C_1 of the next unit.

$$C_1 - C_2 - C_3 - C_4$$

Isopropanol. 2. propanol.

Isopropyl Alcohol. 2-propanol.

Isopropyl Benzene, Cumene, CoH12, CoH5CH (CH3)2. B.P. 152°C Prepared from benzene and benzene in presence of HF.

Isopropyl Acetate, CH₃ COOCH (CH₃)₂. B.P. 88 2°C. Colourless liquid, soluble in organic solvents. Obtained from propylene and hot acetic acid (containing H₂SO₂). Forms azeotrope with 52.5% isopropanol. Used as a solvent.

Isopropyl Chloride, 2-Chloropropane, C3H7C1.

Isopropylidene. The group

Isosbestic Point. A point or points in spectra of transition metals at which extinction co-efficients are equal.

Isostructural. Compounds, substances or elements having the same type of lattice and crystal structure.

Isotherm. A line an a graph joining points of equal temperature,

Isothermal Change. A process that takes place at a constant temperatings rature. The system is always in equilibrium with its surroundings Isothiocyanates. RNCS.

Isotones. Atomic nuclei that have the same number of neutron, but

Isotonic. Solutions having the same osmotic pressure values.

Isotopes. Species of the same element that have same atomic number (similar chemical properties) but different atomic weights, e.g., ¹² C and ¹⁴ C. The isotopes differ in physical properties because of the difference in mass. This property is used for the separation of isotopes. Other methods for isotope diffusion electromagnetic etc.

Isotope Effect. A kinetic isotope effect due to which the rate of a C-H bond breaking is slower when the hydrogen is replaced by deuterium (D) or tritium (T). Less energy is required for

Isotopic Mass (isotopic weight). The mass number of a given

Isotopic Number. The difference between the number of neutrons

Isotopic Weight. See isotopic mass.

Isotropic. A crystalline solid exhibiting identical properties in all directions, e.g., reflection, thermal conductivity, etc. A cube is isotropic.

Isoconic Acid, C₅H₆O₄, H₂C=C-COOH

M,P. 162-164°C. White solid, used as a co-monomer in plastics. IL THE LOCAL CONTRACTOR OF STATE

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e are the west upon a good area from which is the contract of the contract of

J. A measure of coupling constant in nuclear magnetic resonance.

"J-Acid" Dyes. The simple monazo-dyes obtained from "J-acid".

Jalapin, C₃₄H₆₂O₁₈. A glycosidic resin.

Jasper. A hardened siliceous clay used as a gem stone and orna-

Jelutong. A rubber-like material obtained from the tree Dyera Costularia.

J.J. Coupling. See Russel-Saunders coupling.

Jones Reductor. A tube (having zinc amalgam) used to reduce solution prior to estimation.

Joule, J. The SI unit of energy and work and is equal to the work done in moving the point of application of the force of one Newton by one meter in the direction of force, 1 J=1 Nm.

Joules' Law. An apparent law stating that the internal energy of a gas depends only on its temperature and is independent of its pressure and volume. Not applicable at high pressures.

Joule-Thomson Effect. Most of the gases, except hydrogen and their expansion through a nozzle helium, undergo cooling on their expansion through a nozzle under adiabatic conditions. This is known as Joule-Thomson effect. Used for liquefying air, making solid CO₂, etc.

Juniper Tar Oil. Oil of cade.

Juvenile Hormones. Insect hormones used to regulate larval development or reproduction of insects. Used in insect controls.

K

K. Potassium.

- Kainite, MgSO₄ KCl,3H₂O. Occurs in the Stassfurt salt deposits as beds. Forms colourless monoclinic crystals (when pure). Used as a useful fertilizer and a source of potassium salts.
- Kairomone. A material by which an insect is attracted to its food plant.
- Kaolin. Describing certain primary clays, known for their whiteness after calcination and their high refratoriness, e.g. china clay.
- Kaolinite, Al₂ (OH)₃ Si₂O₅. A white clay mineral, apparently amorphous but in reality crystalline (by X-ray analysis). An important constituent of china clay.
- Kapustinskii Equation. For an ionic crystal, the lattice energy (U) is given by the equation:

$$U = \frac{LZ_{+}Z_{-}e^{2}M}{R} \left(1 - \frac{1}{n}\right)$$

where Z_+ =charge on the cation, Z_- =charge on the anion, L=Avogadro's number, e=the electronic charge, R=the equilibrium internuclear distance, M=the Madelung constant, and n=an integer.

The Kapustinskii equation is a modified form of the above equation and is:

$$U = \frac{0.777 \text{ No } \times Z_{+}Z_{-}e^{2}}{(r_{+} + r_{-})}$$

where v= the number of ions per molecule, $r_{+}=$ the thermodynamic radius of caution, $r_{-}=$ the thermodynamic radius of anion, $R=(r_{+}+r_{-})$, the average value of n=9.

Karl Fischer Reagent. A mixture of I2 and SO2 dissolved in pyridine -CH3OH.

Kayser. A proposed unit of frequency 1 cm⁻¹.

Kekule Structure, of Benzene. A structure of benzene with alternate

double bonds, e.g.,
$$\longleftrightarrow$$

Two possible resonating structures contribute to the resonance hybrid of benzene.

Kel-F. A thermoplastic.

Kelvin, K. A thermodynamic temperature unit, zero kelvin (0K) is absolute zero and is equal to 0 of centigrade scale.

Kelvin Scale. See absolute zero.

Kepone, C₁₀H₁₀O. A ketonic solid, used as an insecticide and fungicide.

Keratins. An insoluble protein or group of proteins, belonging to the scleroprotein class, found in the skin, hair, nails, horns, hoofs, etc. Found in α and β form.

Kernite. Na₂B₄O₇, 4H₂O. An important source of borax compounds.

Kerosin. A petroleum fuel, used as lamp oil and with boiling range of 410-570 K.

Ketals. Ketone acetals of general formula

$$R > C < \frac{OR''}{OR'''}$$

colourless liquids obtained by reacting acetylene (C_2H_2) with alcohols in the presence of HgO and BF₃.

Keten, CH₂=C=O. A colourless gas, obtained from acetylene (passing through metallic tubes at 550-800°C). Used as an acetylating agent.

Keto. A group >C=0.

Keto-enol Tautomerism. Describing an equilibrium of organic compounds between two isomers in which one isomer is a ketone and the other an enol. It is due transfer of a hydrogen atom between the oxygen atom of the carbonyl group and an adjacent carbon atom. For example, acetoacetic acid shows tautomerism as shown below:

Ketohexose. A ketose sugar with six carbon atoms and containing ketone group, e.g., Glucose.

Ketols. Organic compounds containing both a keto and an alcohol group. Formed by condensation between two molecules of ketone or by oxidation of Glycols. Exhibit the properties of both ketones and alcohols.

Ketone Bodies, Acetone Bodies. Organic compounds such as propanone (acetone), CH₃COCH₃, acetoacetic acid CH₃COCH₂ COOH, etc.

Ketones. Organic compounds having the CC=O,

group and represented by the general formula $\underset{\mathbf{R}'}{\overset{R}{\longrightarrow}} = 0$,

where R and R' are alkyl groups (may be same or different). They may be aliphatic, alicyclic, aromatic or mixed ketones. In cyclic ketones the carbon of the carbonyl group forms a part of the ring. Formed by passing the vapours of acid over thorium at 400°C or by the dry distillation of Ca or Ba salts. Aromatic ketones are usually prepared by Friedel-Crafts reaction. Aliphatic ketones are generally, liquids whereas aromatic are ketones solids. Soluble in organic solvents such as alcohol and other. Reduced to secondary alcohols and form oximes with hydroxylamine and semi-carbazone with semi-carbazide. Strong oxidizing agents produce a mixture of carboxylic acids. They do not react with Fehling's solution and Tollen's reagents (a distinguishing test from aldehydes). Do not undergo polymerisation.

Ketose. A sugar containing potential keto-(CO) group, e.g., keto-

Ketoximes. Organic compounds containing the group >C=NOH. Formed by the ketone and hydroxyl amine. Reduced to pamines and also undergo Beckmann rearrangement when treated

$$\begin{array}{c}
CH_3 \\
CH_3
\end{array}
C = O + H_2NOH \rightarrow CH_3 \\
CH_3 \\
CO = NOH + H_2O$$
Intensely coloured salts of roll in the sal

Ketyls. Intensely coloured salts of radical anions of ketones, e.g.,

Kevlar. An aromatic amide fibre.

Kieselguhr. A naturally occurring and very porous form of silica-(SiO₂). Used as an adsorbent material and catalyst support.

Kick's Law. A law related to the power consumption in any size reduction process and mathematically it is expressed as :

 $Hp = K \log \frac{D}{d}$

where Hp=power consumed, D=original diameter of, particles, d=final diameter of particle, and K=an arbitrary constant.

Killed Spirits. ZnCl₂ solution used as a flux for solder. Obtained by adding zinc to hydrochloric acid.

Kilo (K). A prefix denoting 103, e.g., 1 kilogram (103 gms.).

Kilowatt-hour, (kwh). A unit of electrical energy, equal to the energy transferred by one kilowatt power per hour. (1 kwh= 3.6×10⁶ J.)

Kinase. An enzyme used to catalyse the transfer of phosphate from ATP to the final substrate.

Kinetics, Chemical. A branch of chemistry used to study the characteristics of chemical reactions, e.g., rate of a reaction, the effect of temperature, pressure and concentration on the rate of a reaction, the mechanism of a chemical reaction, the effect of a catalyst on the reaction, etc.

Kinetic Theory of Gases. A theory of gases with the following assumptions:

- A gas is made up of molecules which are always in a random motion.
- 2. A gas shows a pressure at a temperature.
- 3. The gas molecules collide with each other and with the walls of container.
- 4. The molecular collisions are always perfectly elastic (no loss of momentum of particles). The general kinetic gas equation PV=nRT, and the various properties of gases are derived by applying the law of probability and of particle dynamics to such systems.

King's Yellow. A pigment made up of a mixture of arsenic trisulphide and trioxide.

Kipp's Apparatus. An apparatus for the production of a gas from the reaction of a solid and liquid, e.g., the production of H₂S gas from iron sulphide and dilute sulphuric acid.

Kirchhoff's Equation. An equation derived from the First Law of Thermodynamics co-relating the specific heats of reactants and products in a chemical reaction with the total heat

$$\left(\frac{dQ}{dT}\right)$$
 as represented below:

$$\frac{dQ}{dT} = C_1 - C_2$$

where C_1 and C_2 are the specific heats (heat capacities) of reactnts and products of the chemical reaction.

- Kjeldahl's Flask. A flask used in the estimation of nitrogen by Kjeldahl method.
- Kjeldahl's Method. A method used for the determination of nitrogen in organic compound. The organic compound is converted to ammonium sulphate by boiling with conc. H₂SO₄ in Kjeldahl's flask. The mixture is then made alkaline and ammonia distilled off into standard acid which is determined by volumetric titration methods.
- Knocking. A process responsible for reduced power output which is accompanied by sharp knocking sounds in spark-ignition and compression-ignition engines. It is reduced by adding anti-
- Knock Rating. The tendency of gasoline and other diesel fuels to produce knocking. It is expressed as 'octane number' or
- Kohlrausch Equation. An equation which explains the behaviour of strong electrolytes on dilution and states, that

$$\lambda \infty - \lambda v = k \sqrt{C}$$

where $\lambda \infty$ = the equivalent conductivity at infinite dilution, λv =the equivalent conductivity at volume v, k=a constant, and C=the concentration of electrolyte. The equation holds

Kolbe Electrolysis. A process used for the preparation of saturated and unsaturated hydrocarbons by the electrolysis of alkali salts of aliphatic carboxylic acids, e.g., ethane from concentrated solution of sodium or potassium acetate.

At anodé: 2CH3COO->CH3-CH3+2CO2+2e-

At cathode: $2H_2O + 2e^- \rightarrow 2OH^- + H_2$

This is a coupling reaction and only the alkanes with even number of carbon atoms in the chain can be prepared by this

Kr. Krypton.

Kreb's Cycle, Citric Acid Cycle. A cycle of reactions in cell metabolism to achieve the controlled oxidative breakdown of acetyl coenzyme A, derived from carbohydrates and fatty acids. Other molecules (e.g., amino acids) can be altered so that they also can be used in the cycle. The various products of the cycle are CO₂ and reduced co-enzymes such as CoASH, NADH, NADPH, FADH₂, which serve as energy sources, under aerobic conditions. This occurs through the respiratory chain

Krilium. The trade name for a soil conditioner.

Kroll Process. A method used for the production of some metals by reduction of metal chloride with magnesium, e.g., the production of Ti from TiCl,.

TiCl₄+2Mg→2MgCl₅+Ti

Krypton, Kr., At. No. 36, At. wt. 83.80, M.P.-157.3°C, B.P.-153.4°C. D 3.73. A colourless, monoatomic, inert gas belonging to zero group of periodic table. Electronic configuration $4s^2 4p^6$. Obtained by fractional distillation of liquid air. Used in fluorescent lights and in photographic flash lights.

Kurchatovium. 104th element in periodic table and belongs to post-actinide series.

Kurrol Salt, NaPO3. A long chain metaphosphate.

Kynurenic Acid, C10H7O3N. 4-hydroxyquinoline-2-carboxylic acid.

Kynurenine, C10H12N2O3. An intermediate substance formed in the m etabolic breakdown of tryptophan.



La. Lanthanum.

Label. A stable or radioactive nuclei used for the investigation of some processes, e.g., a chemical reaction. Radioactive isotopes are used as 'label'.

Labile Complex. A complex which takes part in very fast reactions. mainly ligand exchange reactions, e.g., [Co (H₂O)_n]²⁺.

Laccase. An enzyme found in various bacteria, in mushrooms, in higher plants and fruits and which is responsible for the oxidation of polyphenols to the respective quinones.

Lactalbumin. A protein of albumin class with M. wt. of about 175000. Present in milk.

Lactams. Organic compounds containing the -NHCO- group as part of a ring in the molecule. Amino acids when heated lose part of a ring in the flow (a reaction of the amino group with water to form factains (a molecule). In general, they are caboxyl group within the molecule). Lactams may be ring compounds colourless poisonous solids. Lactams may be ring compounds containing five, six and eight atoms.

β-Lactic Acids. Hydracrylic acid.

Lactic Acid, 2-hydroxypropionic Acid, C₃H₆O₃, CH₃.CHOH.COOH. D-lactic acid, M.P. 18°C. L-lactic acid and D-lactic acid are known. L-lactic acid (sarcolactic acid), occurs in muscles due to the breakdown of carbohydrate (M.P. 25-26°C). Lactic acid is manufactured by the fermentation of pure sugars or sugar containing materials, e.g., starch hydrolysates.

$$C_{12}H_{22}O_{11}+H_2O\rightarrow C_6H_{12}O_6+C_6H_{12}O_6$$

$$C_6H_{12}O_6- \xrightarrow{\text{BAL}} C_6H_{12}O_6 - \xrightarrow{\text{CHOH-COOH}}$$

$$C_6H_{12}O_6 - \xrightarrow{\text{(Bacillus acidii lactii)}} 2CH_3-CHOH-COOH$$

Used as a mordant for dyeing wol, as acidulant in candles, in medicines and in plastic industry.

Lactide, C₆H₈O₄. Occurs in optically active forms. Prepared by the slow distillation of concentrated solutions of the lactic acids.

Lactides. Compounds containing the group and obtained from CO-O-CH- α-hydroxy fatty acid by the loss of water —CH-O-CO

Lactones. Anhydrides formed by elimination of water between the hydroxyl and carboxyl groups of hydroxyacids, e.g., lactones

$$\begin{array}{c} \text{O} \\ \text{CH}_2-\text{C} \\ \text{and } \text{H}_2\text{C} \\ \text{O} \\ \text{CH}_2-\text{CH}_2 \\ \text{\delta-valerolactone} \end{array}$$

Lactore, C₁₂H₂₂O₁₁ (milk sugar). Found in milk of all animals (6% in human milk, 4% in cow's milk). Optically active, found in two series of isomers and composed of glucose and gallactose unit.

Laevo Rotatory. See optical activity.

Laevulinic Acid, Laevulic Acid, C₅H₈O₃, CH₃COCH₂CH₂COOH. M.P. 33-35°C. Obtained from cane sugar or starch and con. HCl. Used in cotton printing.

Lake Asphalt. A naturally-occurring mixture of bitumen with universal and organic matter.

Lake. A pigment obtained from organic dyestuffs and inorganic compounds.

Lambda (A) Print. The transition temperature of He1 and He11.

Lambert's Law. The law states that layers of equal thickness of a homogeneous material absorb equal proportions of light, or mathematically, $I = I_0 e^{-kd}$

where I=the intensity of the transmitted light, I_0 =the intensity of incident light, d=the thickness of the layer, and k=a constant (absorption co-efficient), characteristic of the substance. k depends on the wavelength (λ) of light.

Lameller Compounds. Compounds with a crystal structure made up of thin plates or layers, e.g., talc [Mg₃(OH)₂Si₄O₁₀] and pyrophyllite [Al₂(OH)₂Si₄O₁₀].

Laminarin (C₆H₁₀O₅)_n. A white powder, obtained from β-D-glucopyranose units.

Laminates. Laminated sheet materials prepared by bonding layers of reinforcing agents such as cotton cloth, paper, asbestos fabric, and glass clothes by synthetic resins. Amino resins or phenol resins can also be used as binding materials for sheets. They have high track resistance, high water and acid resistant power and good tensile strengths. Used for constructing gears, bearings and as decorators.

Lamp Black (a black pigment). A finely divided form of carbon obtained by the incomplete combustion of an organic compound.

Langmuir. A unit for absorption. 10⁻⁶ torr adsorbent exposed to

Langmair Adsorption Isotherm. A theoretical equation given by:

$$\theta = \frac{bp}{1 + bp}$$

where θ=the fraction of the surface covered by a gas adsorbed at a plane solid surface, p = the pressure of the gas in equilibrium with the solid surface, and b a constant called the absorption co-efficient, an equilibrium constant for the absorption

(A monolayer formation of gas is assumed on the solid

Lanoceric Acid, C₃₀H₆₀O₄. A saturated dihydroxy acid present in wool. (M.P. 105°C).

Lanolin. M.P. 37°C. A pale yellow substance obtained from wool.

Lanthanides. A group of elements (from Lanthanum to Lutetium) whose 4f orbitals are being filled. A closely related series of 15 elements having electronic configuration $4f^{x}6s^{2}$ (excluding La) and Gadolinium and Lutetium have an additional 5d electron. The La itself has no f-electrons [(Xe)5d¹6s²] but

All metals are shiny with a characteristic oxidation state of M³⁺ and with similar chemical properties. The elements are separated by chromatography or ion-exchange.

Lanthanide Contraction. A regular decrease in ionic and atomic

Lanthanum, La. At. No. 57, At. wt. 138'9055. M.P. 920°C, B.P. 3454°C, D 6'14. A soft, ductile, malleable silvery element belonging to the lanthanide series. longing to the lanthanide series. A paramagnetic element found in many minerals such as monazite and bastnasite. Used in alloys, as a catalyst, in carbon-arc search lights, etc.

Lapis Lazuli (ultramarine, mineral). A semi-precious stone.

Laser. A device to produce beams of monochromatic light of very great intensity where the waves are coherent (the wave fronts are in steps with one another). Example of a solid laser is the ruby crystal (Al₂O₃ containing about 0.5% Cr). Used in devices where controlled energy is required, in Raman

Lassaigne's Test. A test for the detection of nitrogen, halogen or sulphur in an organic compound. The organic compound is converted into the corresponding salt of nitrogen, halogen or

sulphur by fusion with metallic sodium. Nitrogen is detected by the formation of a Prussian blue colour with ferrous sulphate solution (containing FeCl₃) and HCl. Halogens in the compound are detected by AgNO₃ solution to form precipitates of silver halides. Sulphur is detected by sodium nitroprusside.

Latent Heat (a) of fusion-The heat absorbed when a solid is converted to liquid at the melting point.

(b) of vaporization-The heat absorbed when a liquid is converted to a gas at the boiling point. Latent heats are generally expressed in J mol-1.

Latent Image. A term used in photography.

Latex. An emulsion or suspension containing rubber particles. Many natural rubber latex were obtained from the bark of some trees. Many synthetic rubbers and polymers are produced as latex, e.g., PVC. Used for the direct manufacture of rubber, adenine and for paints.

Lattice. A regular three dimensional arrangement of points in a crystal. The point may contain the particles such as atoms, ions or molecules in a solid. Lattice structure can be studied by X-ray diffraction method.

Lattic Energy. A measure of the stability of an ionic solid structure with respect to ions in the gas. It can be defined as the energy required to break down 1 mole of an ionic lattice into its constituent ion in the gaseous state and separated from each other to an infinite distance or the energy released when ions of opposite charge are brought together from infinity to form one mole of a given substance.

 M^+X^n (solid) $\rightarrow M^+(g) + nX^-(g)$

It can be calculated by Kapustiniskii equation.

Laue Pattern. Patterns obtained on photographic plates during the X-ray investigation of crystal structures.

Laughing Gas. Dinitrogen monoxide.

Lauric Acid, C₁₂H₂₄O₂, n-Dodecyl Acid. M.P. 44°C, needles. A fatty acid occurring in milk, laurel oil, coconut oil, palm oil, etc.

Laurite, (Ru, Os)S2. A source of Ru and Os.

Lauryl Alcohol, Dodecyl Alcohol, C12H26O. An alcohol, M.P. 24°C, insoluble in water but soluble in alcohol. Manufactured by reduction of ethyl laurate or glyceryl trilaurate by hydrogen under pressure. Used in the manufacture of detergents.

Law of Constancy of Angles. See interfacial angles.

Law of Isomorphism. Mitscherlich's Law.

Law of Mass Action. At constant temperature, the rate with which reactants take part in a chemical reaction is directly proportional to their active masses. For the reaction

aA+bB→Products

the rate of the reaction ∞[A]2 $\alpha[B]^b$

 $=k[A]^a[B]^b$

where [A] and [B]=the active masses of reactants, a and b the number of moles of reactants and k=the rate constant of the

Active mass=Activity coefficient × Concentration

For dilute solutions, activity coefficient is unity and, hence, the active mass is numerically equal to the molar concentration.

Law of Octaves. See Newland's Law.

Lawrencium, Lr. At. No. 102. A radioactive transuranic element of the actinoid series, does not occur in nature. Several shortlived isotopes have been synthesised by bombardment of Cf targets with 10B and 11B nuclei in a cyclotron or linear accelerator. Known only in +3 oxidation state. Does not form any

LCAO Method. A method used to calculate molecular orbitals which is based upon the concept that the molecular orbitals can be expressed as a 'linear combination of atomic orbitals'.

Leaching. A process of washing out of a soluble material from an

Lead, Pb. At. No. 82, At. wt. 207'2 (approx.), M.P. 327'502°C, B.P. 1740°C, D 11'34. A dense, dull grey and soft element of group IV of periodic table, occurs as galena (PbS), anglesite (PbSO₃), litharge (PbO) and cerussite (PbCO₃). Obtained from galena by concentrating (by froth flotation method) roasting

 $2PbS + 3O_2 \rightarrow 2PbO + 2SO_2$ PbS+2O₂→PbSO₄ $2PbO + 3C \rightarrow 2Pb + 2CO$ $2PbO + PbS \rightarrow 2Pb + 2SO_2$ $PbSO_4 + 2C \rightarrow Pb + 2CO + SO_2$

(Silver is, generally, recovered from crude lead). Lead is a mixture of its various isotopes. It has outermost configuration s^2p^2 and exhibits similar properties for two types of lead, e.g., Pb(II) and Pb (IV). Forms mixed oxides such as Pb (II) O (lead and Pb3O4.

(plumbus and plumbic) have low M.Pts. Used extensively in alloys, in storage batteries, for covering electrical cables, etc. Also used in the manufacture of lead accumulators, red lead. and white lead.

Lead Accumulator. An electrical accumulator made up of spongy lead plates connected in series to the negative terminal and lead oxide plates connected to the positive terminal. The plates are interleafed and the electrolyte is dilute sulphuric acid. When a current is passed through the accumulator PbO2 is formed on one sets of plates and Pb on the other, and some H₂SO₄ is formed. When electricity is generated in the accumulator, H₂SO₄ acts on Pb, PbO and PbO₂ to produce PbSO₄ and water. The emf when fully charged is about 2.2 volts. To recharge the accumulator, electric current is passed through it in a direction opposite to that of the current supply.

Lead Azide, Pb(N₃)₂. Used in explosives.

Lead Bromide, PbBr₂. Soluble in hot water.

Lead Carbonate, PbCO₃. A white poisonous powder occurring in cerussite. Forms basic carbonates, e.g., Pb₃(OH)₂(CO₃)₂, from lead; used in paints.

Lead Carbonate Hydroxide (white lead), 2PbCO3, (PbOH)2. See

Lead Chromates, PbCrO₄. Precipitated from solution of Pb²⁺ and CrO42- or Cr2O72-.

Lead Fluorides,

PbF₂-Obtained from aqueous solutions.

PbF₄-A strong fluorinating agent.

Lead Nitrate, Pb(NO₃)₂. Used in calico printings, as a mordant in dyeing and for the manufacture of yellow pigment.

Lead Organic Derivatives. Derivatives of lead (II), e.g., tetramethyl lead [(C₄H₅)₄Pb] which are anti-knock additives for petroleum.

Lead Oxides

Lead (II) Oxide, PbO. A yellow crystalline powder formed by roasting molten lead in air. Litharge is obtained by heating PbO above its melting point while massicot is obtained below its melting point. Used in the manufacture of paints and var-

Lead (IV) Oxide, PbO₂ (lead dioxide). A dark brown solid prepared electrolytically or by reducing lead oxide with KClO₃. Used in the manufacture of matches.

154 Lead Silicates

Lead Silicates. Silicates of lead insoluble in dilute acids and used in glazing pottery and in glass manufacture.

- Lead Sulphate, PbSO₄. A white solid occurring in the mineral anglesite. Insoluble in water, forms basic lead (II) sulphate when shaken together with lead (II) hydroxide and water. Used
- Lead Sulphide, PbS. A black solid occurring in galena. Used as a rectifier in electrical components.
- Lead Tetraethyl (tetraethyl lead), Pb (CH₂CH₃)₄, TEL. A poisonous liquid, insoluble in water and manufactured by the reaction of an alloy of sodium and lead with 1 chloroethane followed by steam distillation. Used as an additive in internal combustion engine to increase the octane number of fuel.

Leaf Filter. A device of filtration.

Lean Gas. A fuel gas with a calorific value of 100-450 BTU/ft3.

Leather. Animals and other skins which are tanned.

Leblanc Process. An obsolete process for the manufacture of

Le Chatelier's Principle. If a system is in equilibrium, a change in any of the factors (e.g., P, T or concentration) that determine the condition of equilibrium, will cause the equilibrium to shift in such a way as to minimize the effect of this change.

$$N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$$

increasing the pressure will cause the equilibrium to shift towards

- Leclanche Cell. A primary voltaic cell made up of a carbon rod anode, a Zn cathode and 10-20% solution of ammonium chloride as an electrolyte (wet cell). MnO2 and crushed carbon surround the anode (as a depolarizer). The dry cell is used for torch batteries, transistor radios, etc. It is made up of a mixture of NH₄Cl, ZnCl₂, flour and gum which acts as an electrolyte
- LEED. A technique for investigating the two dimensional surface structure of solids. The surface to be investigated is bombarded with electrons of low energy (between 6-600 volts), and the electrons diffracted by surface atoms of the solid are collected on a fluorescent screen. Used to study surface structures before and after chemisorption or any other chemical reaction

Legumelin. A protein of albumin class.

Legumin. A protein of globulin class.

Lel. A term used in chelate chemistry.

Lepidolite, Si₃O₉ Al₂ (Li, K)₂ (F, OH)₂. A lithium potassium mica. Occurs chiefly in pegmatites. Used as a raw material for the manufacture of Li.

Leucine, a-Aminoisocaproic Acid, C6H13O2N,

Soluble in water up to 2%. One of the important amino acids obtained from proteins.

Leucite, K₂Al₂Si₄O₁₂. Potassium aluminosilicate; occurs as a primary constituent of volcano rocks.

Leucopteron, C₆H₆O₃N₅. A white pigment prepared from the wings of butterflies.

Leucosin. A protein of albumin class. Present in the seeds of wheat, rye and barley.

Leuco Compounds. Colourless compounds obtained from dye stuffs.

Leven, $(C_6H_{10}O_5)x$. A water-soluble polymer of fructose present in some grasses.

Lewis Acid. A substance (molecule, ion or radical) that can accept a pair of electrons to form a coordinate bond. Acid is thus an electron-pair acceptor and base is a donor of electron pair. For example, the reaction Et₃N+BCl₃→Et₃N: BCl₂ is an acid base neutralization reaction, though, there is no proton nor any OH group (in chlorobenzene medium).

$$H^++: S: H^- \longrightarrow H: S: H$$

Metal ions in coordinate compounds are also Lewis acids as they also accept electron-pair.

Lewis Base. An electron-pair donor to a Lewis acid.

Lewisite, C₂H₂AsCl₃. A war gas poison. Destroyed by alkalis and oxidizing agents.

Li. Lithium.

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Libration. The rotational oscillations of the whole molecule within the lattice are known as libration modes.

Licanic Acid, C₁₈H₂₈O₃.

- **Liebermann's Reaction.** A test for the identification of $-N \cup$ or -OH groups.
 - -NO group-Concentrated sulphuric acid+substance+a crystal of phenol, warm-a green colour develops which changes to red in water and black blue with alkalis.
 - -OH group- Substance+a crystal of NaNO₃+conc. H₂SO₄, warm, dilute and add alkali—the formation of a distinct colour.
- Ligand. A molecule or ion which forms a coordinate bond to a metal atom or ion in a complex, e.g., NH₃ is a ligand in [CO (NH₃)₆]Cl₃.
- Ligand Field Theory. See crystal field theory.
- Light (visible radiation). An electromagnetic radiation which is detected by human eye. The wavelength (λ) range is between 400 nm (far 'red') and 700 nm (far 'violet').
- Light-Heavy Selectivity. A phenomena in solvent extraction in which the extraction power of a solvent depends on the M. wt. of the component extracted.
- Light Petroleum. The refined fraction of petroleum or shale oil boiling from 40°-60°C. Contains mainly the pentanes and hexanes. Used as a solvent for the fats and waxes.
- Light Scattering. A phenomenon of scattering of light by suspended particles (size of the particles should be slightly smaller than molecular weights of macromolecules and to detect the end point in precipitation titrations.
- Light Stabilizers. See ultra-violet absorbers.
- Lignan. A group of natural products obtained, generally by etheral or alcoholic extraction, from the wood or exuded resin of the coniferae and other plants and characterised by the presence of a 2, 3-dibenzylbutane skeleton in the molecule
- Lignin. An amorphous, highly polymeric substance occurring in cellulose in lignified tissues. Dissolved out from pulp by SO₂ and lime water. Responsible for the strength of wood (contains 25-30%). Used as a filter for plastics, as emulsifier, etc.
- Lignite (brown coal). A type of soft fuels, similar to the peats (containing large quantities of water). Occurs in seams quite

- Lignoceric Acid, C₂₄H₄₈O₂. A fatty acid occurring in free and combined state in several oils, fats and waxes.
- Ligroin, Petroleum Naphtha. A fraction of petroleum boiling between 90°-120°C. Contains chiefly heptanes and octanes. Used as a solvent.
- Lime. A white solid formed by heating calcium in oxygen or by the thermal decomposition of CaCO₃. Manufactured by heating limestone at 550°C.

 $CaCO_3 \rightarrow CaO + CO_2$ Lime stone

Used for the formation of slags and in white washings.

Lime Stone. A natural variety of calcium carbonate (calcium carbonate in the form of calcite). Commercial lime stone contains iron oxide, magnesia, alumina, silica and sulphur with CaO and MgO. Used for making calcium compounds, carbondioxide and cement.

Lime Water (calcium hydroxide), Ca(OH)₂ (slaked lime). See calcium hydroxide.

Limiting Density (of a gas). If a mass w gm of a gas occupies v litres at 0°C and pressure p atmospheres, the quotient w/pv is the density of the gas per unit pressure. But gases do not obey Boyles' Law strictly. $\frac{w}{}$ varies with p. The value of this

Boyles' Law strictly, $\frac{n}{pv}$ varies with p. The value of this quotient when p approaches zero is known as the limiting density of the gas. Alternatively, it is the density of an 'ideal' gas.

Limonene, C₁₀H₁₆. An optically active terpene. Obtained by the action of heat on limonene. B.P. 176°C.

Linalool, C₁₀H₁₈O. B.P. 198-199°C. Present in many essential oils.

Optically active. CH₃ OH

158 Linamarine

Linamarine, C₁₀H₁₇O₆N (acetonecyanhydrin-β-glucoside). C₆H₁₁O₅.
O. (CH₃)₂. CN. M.P. 143°-145°C. Occurs in young flax plants and Phaseolus lunatus. Used as a primary material for rubber synthesis.

Lindane. The Y-isomer of BHG, HCH.

Linde Process. A process for liquefying O₂ and N₂ by compression to about 200 bar (20MN/m²) followed by refrigeration and fractionation.

Line Spectrum. A spectrum consisting of a number of discrete lines corresponding to single wavelength of emitted or absorbed radiation (also single frequency). Each line on spectrum (at a frequency) corresponds to the appropriate electronic transition. It is a characteristic of an element in the atomic state.

Linkage. Bond.

Linoleic Acid, Linolic Acid, C₁₈H₃₂O₂. A liquid, B.P. 229-230°C/16 mm. Unsaturated fatty acid occurring in linseed oil, cotton seed oil and several other vegetable oils. Used to treat animals which fail to lay down fat and suffer from skin and kidney diseases.

Linolenic Acid, $C_{18}H_{30}O_2$. A fatty acid occurs in linseed oil. Used as an essential fatty acid.

Linseed Oil. Linolenic acid—containing oil of vegetable origin.

Used as a drying oil, in resins and margarine.

Lipases. A class of hydrolases (enzymes).

Lipids. They are fatty acids which are soluble in organic solvents and insoluble in water. Classified as simple lipids, fats and waxes; compound lipids, the phospholipids and cerebrosides; and derived lipids, the fatty acids, sterols, and alcohols.

Liquid. The state of matter in which the constituent particles are loosely bound by intermolecular forces. Liquids can change their shape with a fixed volume.

Liquid Air. A nale blue liquid and a mixture of liquid oxygen and liquid nitrogen.

Liquid Crystals. A cloudy liquid that has definite ordered structure. Obtained by heating a substance (e.g., ammonium oleate), higher temperature. Each transition takes place at a fixed dimensional order (cholesteric or smetic phases). Those who which produce liquid crystals are made up of molecules which

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have high degree of asymmetry (long thin molecules or flat planer molecules) and which allow only a little rotation in the liquid state at low temperature. Used in thermometer units, in spectroscopy, etc.

- Liquid Extraction. A process of extracting a liquid component from a liquid mixture by a suitable solvent.
- Liquid Oxygen Explosives, LOX Explosives. Explosives of liquid oxygen and a fuel (carbon black).
- Liquid Paraffin. A highly refined white oil used as a laxative.
- Liquids, Structure of. A liquid has an order intermediate between the perfect disorder of molecules in a gas and a three dimensional order in a solid (no range order).
- Liquidus Curve. If the freezing points of the mixture are potted as a graph using temperature and composition as ordinates (T-X diagram), the line joining the freezing points is known as called a liquidus curve. If the temperature at which the last traces of liquid just solidify are plotted against composition, the resulting curve is called a solidus curve.
- Liquified Natural Gas (LNG). Natural gas liquified by cooling to about -160°C and at 1 atmosphere pressure for transportation purpose.
- Liquified Petroleum Gas (LPG). A domestic and industrial fuel and a mixture of C₃ and C₄ hydrocarbons (e.g., commercial butane and propane). Stored and transported as liquids. Calorific value 10⁵ kJ/m³.

Litharge, PbO. (Lead oxide).

Lithia, Li2O. Lithium oxide.

Lithium, Li. At. No. 3, At. wt. 6.940, M.P. 180°C, B.P. 1336°C, D 0.54. Occurs as petalite (LiNaH)Al(Si₂O₅)₂. A light silvery moderately reactive metal, the first member of the alkali metal group (I group) of periodic table. Chief minerals of lithium are triphylite [(LiNa)₃PO₄+(Fe, Mn)₃(PO₄)₂], lepodolite or lithium mica. (Li, K)₂ Al₂(F, OH)₂Si₃O₉], amblygonite [LiAl(F, OH)PO₄] and spodumene [Li Al(SiO₃)₂], obtained by electrolysis of a fused mixture of lithium and potassium chlorides, with a C anode and an iron cathode. Solutions of Li salts in solvents such as pyridine, acetone, etc. On electrolysis produce pure Li. Forms LiH, Li₂O, Li₃N when reacted with hydrogen, oxygen and nitrogen. Li compounds impart a characteristic purple colour to flames. Li compounds are used in organic synthesis (alkyl and aryl compounds), in ceramics and glasses (LiOH), in air conditioning plants (LiCl and LiBr), in food stuffs, in fungicides, etc.

- Lithium Aluminium Hydride. LiAlH₄, A reducing agent; used in the formation of aluminium hydrides.
- Lithium Carbonate, Li₂CO₃. A white solid obtained from a lithium salt and sodium carbonate solution. Sparingly soluble in water (distinction from other I group carbonates). Resembles the carbonates of Mg and Ca (diagonal relationship).
- Lithium Chloride, LiCl. A white solid obtained from Li₂CO₃ or Li₂O in HCl solution. Forms dihydrate [LiCl. 2H₂O] below 19°C and becomes anhydrous at 93.5°C. Dissolves in organic solvents. Forms cubic crystals. A most deliquescent substance which forms mono-, di- and tri-hydrates.
- Lithium Hydride, LiH. A white crystalline solid formed from Li and H₂ at 500°C. Reacts with water to form Li(OH) and H₂. Used as a reducing agent for the preparation of other hydrides.
- Lithium Hydroxide, LiOH, H₂O. A strong base, obtained from a Li salt or ore and Ca(OH)₂. Used in greases, storage batteries and for absorbing CO₂.
- Lithium Organic Derivatives. Prepared by RCl in C₆H₆ or petroleum ether and Li (where R is an alkyl group). Generally polymeric, e.g., (LiMe)₄. Used in industrial and laboratory synthesis and in alkene-polymerization.
- Lithium Oxide, Li₂O. A white solid obtained from Li and O₂. Forms LiOH with water.
- Lithography. A printing process in which the printing plate is generally made of grained Al or Zn, which is coated with the light sensitive materials.
- Lithopone, Charlton White, Orr's White. A mixture of barium sulphate (70%) and zinc sulphide (30%) by weight. Used as a
- Litmus. A colouring matter obtained from lichens by oxidation in presence, of ammonia. Available in blue cubes which are dissolved in water to act as an indicator. The pH range is 4.5 to acid).
- Litre, l. A unit of volume (10⁻³ m³) equal to 1600 cm³.
- LNG. Liquefied natural gas.
- Lone Pair. A pair of electrons which does not take part in bonding, e.g., ammonia has a lone pair of electrons (:NH₃). Lone
- Long Period. See period.

Loschmidt's Number. The number of molecules of an ideal gas in a unit volume at standard conditions of temperature and pressure; equal to 2.687×10^{19} cm⁻³.

Low Energy Electron Diffraction. LEED.

Lowering of Vapour Pressure. A colligative property of solutions in which the vapour pressure of the solvent is lowered by the addition of a non-volatile solute. The magnitude of lowering of vapour pressure is directly proportional to the number of particles of solute in the solution.

Low-spin State. The minimum numbers of unpaired electrons.

Low Temperature Baths. Baths used for freezing and distilling gases, e.g., liquid $N_2(-195^{\circ}C)$, liquid $O_2(-183^{\circ}C)$, solid $CO_2(-78.5^{\circ}C)$, isopentane $(-140^{\circ}C)$, pentane $(-126^{\circ}C)$, C_6H_5 $Cl(-23^{\circ}C)$, etc.

Lox Explosives. Liquid oxygen explosives.

LTC. Low temperature carbonization.

Lu. Lutetium.

Lubricating Greases. Solid or semi-fluid lubricants obtained by mixing a liquid lubricant and a thickening agent, e.g., greases containing soaps of Li, Ca, Ma, Al, etc. (soapy greases). The base lubricant is usually petroleum oil whereas thickener is usually a soap or a soap mixture. Used as lubricants at high temperatures (non-soapy greases containing finely divided solids), as Bentone greases (organo-clays).

Lubricating Oils. Mineral oils derived from crude petroleum, vegetable oils (used as 'compounded' oils) and synthetic fluids (used for aviation turbines) which are used for the purpose of lubrication. For example, esters, polyglycols, silicones and halogenated hydrocarbons.

Luciferins. Substances that form bioluminescence.

Luminous Paints. Compounded from solid material (such as CaS) which are suitably processed by the addition of traces of heavy metals, can be made to show phosphorescence.

Luminescence. The particles (electrons) in an atom are excited to higher energy states. They then return to lower energy states with the emission of electromagnetic radiation. If the lumini, scence persists even after the source of excitation is removedit is known as phosphorescence, if not, it is called fluorescence.

Lumisterol. Vitamin-D.

- Lurgi Coal Gasification Process. The gasification of coal under pressure with mixtures of steam and O2: A method for the production of CO and Ha.
- Lutein, Luteol, Xanthophyll, C₄₀H₅₆O₂. A pigment with the normal carotenoid structure. M.P. 193°C, soluble in organic solvents.
- Lutetium, Lee. At. No. 71, At. wt. 174.96, M.P. 1656°C, B.P. 3315°C. D 9.84. A silvery element of the lanthanide series (4f series). Occurs in association with other lanthanides. Nuclides emit pure β radiations. Used in cracking polymerization and as a catalyst. Forms typical compounds in +3 state.
- Lutidine, C7H9N. An oily liquid (dimethyl pyridine).
- Lux, lx. An SI unit of illumination and is equal to the illumination produced by a luminous flux of one lumen falling on a surface of 1m2. (1 lx=1 lmm-2).
- Lycopene, C₄₀H₅₆. M.P. 175°C. A red pigment of tomatoes, rose
- Lyman Series. A series of lines (discovered by Lyman) in the ultraviolet spectrum emitted by excited hydrogen atoms. The wave length (λ) of the radiation in the Lyman series is given by

$$\frac{1}{\lambda} = R \left(\frac{1}{12} - \frac{1}{n^2} \right)$$

where n is an integer and R is the Rydberg constant.

- Lyophilic. Sols in which the dispersed particles are solvent loving
- Lyophobic. Sols in which the solvent particles are unsolvated and are easily coagulated (solvent hating).
- Lyotropic Series. A term applied to colloidal solutions and can be defined as the arrangement of different anions (of salt) in order of their salting-out efficiency of colloidal sols.

Lysergic Acid, C₁₆H₁₆N₂O₂. M.P. 238°C.

Lysine, a-e-Diaminocaproic Acid, C6H14O2N2 CH₂—(CH₂)₃—CH—CH₂COOH | | NH₂ NH₂

Colourless crystals, M.P. 221°C, soluble in water and naturally oscurring acid is dextrorotatory $[\alpha]$ $p^{20} + 14.6^{\circ}$. An essential amino acid obtained from protein hydrolysis.

- Lysol. A soapy solution containing 50% isomeric cresols. Dilute solutions are used as disinfectant and antiseptics in surgery.
- Lysosomes. An enzyme which destroys many cell constituents.

Lysolecithin. A substance obtained from enzyme lecithase of cobra venom on lecithin.

Lysezyme. An enzyme with the ability to destroy certain bacteria. Present in nasal mucosa, eggwhite and in various animal tissues.

Lyxose, C5H10O5. A synthesised pentose sugar.

M

Macleod's Equation

$\gamma = K(D-d)^4$

where γ is the surface tension, D is the density of liquid and d is the density of vapour at the same temperature as that of the liquid.

Macrolides. Compounds having a macrocyclic ring and show antibiotic activity. Active against gram-positive bacteria.

Macromolecule. A large molecule with a high M. wt. (more than 10,000), e.g., polymers and proteins.

Macromolecular Crystal. A crystal made of atoms joined together by covalent bonds resulting in the formation of giant three dimensional or two dimensional network, e.g., diamond.

Maddrell Salt, NaPO₃. A long chain phosphate obtained from Na₂H₂P₄O₇ at 230—300°C.

Magnalium. A group of aluminium-magnesium alloys.

Magnesia. See magnesium oxide.

Magnesite, MgCO₃. A native form of magnesium carbonate used as a source of Mg-compounds and in the manufacture of refractories.

Magnesium, Mg. At. No. 12, At. wt. 24'305, M.P. 648'8°C, B.P. 1090°C, D 1'74. Electronic configuration of outermost orbit is 35°2. A light metallic element belonging to II group of periodic table (the alkaline earth). Occurs in minerals such as brucite [Mg(OH)₂], carnallite (MgCl₂.KCl.6H₂O), epsomite (MgSO₄. 7H₂O), magnesite (MgCO₃), dolomite (MgCO₃.CaCO₃) and

also MgCl2 in sea water. Obtained by electrolysis of MgCl2halide melt or by reduction of fused dolomite with ferro-silicon. Burns in air after ignition. Used in glass, ceramics, fillers (MgCO₃ MgCl₂), catalysts, medicinal uses, refractories, sugar refining, cement, paper manufacture, in tanning, etc. Also used in light weight alloys (with Zn and Al). Forms co-ordination compounds (ionic nature of Mg). Organometallic compounds of Mg are used in synthetic organic chemistry. Forms Grignard reagents, RMgX and related dialkyls, R₂MgX.Mg-complexes are used in photosynthesis in plants (chlorophyll).

- Magnesium Alloys. Light alloys used in aircraft industry and for
- Magnesium Carbonate, MgCO₃. A white solid that occurs in magnesite and in dolomite. Sparingly soluble in water and is used as an antacid (basic carbonates-3MgCO₃, Mg(OH)₂, 4H₂O). Heating the solution to 50°, forms MgCO₃, 3H₂O. Obtained from CO₃²⁻ and Mg²⁺ solutions which dissolves in excess CO₂
- Magnesium Chloride, MgCl₂. A solid that exists in many hydrated forms, most commonly as the hexahydrate MgCl₂.6H₂O. On heating, this hydrated forms gets hydrolysed by its water of crystallization, with the formation of HCl(gas) and an oxide:

$MgCl_2+H_2O\rightarrow MgO+HCl$

anhydrous chloride is prepared by evaporating an aqueous solution in an atmosphere of hydrogen chloride. Used in cotton industry, in cement, etc. Fused MgCl₂ is electrolysed to give Mg and Cl₂ on a commercial scale. Forms complexes.

Magnesium Hydrogen Carbonate, Mg(HCO₃). A solid at room temperature. Obtained from water containing CO2 and mag-

An important cause of temporary hardness.

- Magnesium Hydroxide, Mg (OH)₂. A white solid sparingly soluble in water. Obtained from an alkali and a soluble Mg compound [$(Mg^{2+}+2OH^{-}\rightarrow Mg\ (OH)_{2}]$]. Occurs as brucite. Used as an
- Magnesium Nitrate, Mg (NO₃)₂. A stable salt forms 9, 6 and 2 hydrates, obtained by heating hydrates with HNO3.
- Magnesium Nitride, Mg₃N₂ (a nitrogen remover). Obtained from Mg and nitrogen above 300°C. Hydrolysed to NH₃ and
- Magnesium Oxide, MgO (magnesia). A white solid occurring in mineral periclase. Prepared by heating Mg in O2. A basic oxide used as refractory lining for metal, glass and cement furnace.

Magnesium Peroxide, Mg₂O₂. A white insoluble oxide obtained from sodium or barium peroxide and a concentrated solution of Mg salt. Used as a bleach for dyestuffs and silks.

Magnesium Phosphate, Mg₃(PO₄)₂, 8H₂O. Obtained from aqueous solutions. Used in estimation of Mg²⁺ and (PO₄)³⁻.

Magnesium Silicates. Used as an antacid.

Magnesium Sulphate, MgSO₄. Obtained from MgCO₃ and H₂SO₄ as MgSO₄, 7H₂O (Epsom salt). Occurs in kieserite or reichardite. Magnesium sulphate is used as a purgative drug and as an antidote of Ba and barbiturate poisoning.

Magneson, p-nitrobenzeneazores orcinol, C₁₂H₉N₃O₄. A brownish-red powder, soluble in NaOH. Used for the detection and estimation of Mg (a blue lake in alkaline solution).

Magnetic Moment. For a paramagnetic substance

$$\mu = 2.84 \sqrt{\chi_m} \sqrt{(T-\theta)}$$

where μ =magnetic moment, χ =magnetic susceptibility, χ_m = molar susceptibility, T=the absolute temperature, and θ =a constant (curie temperature).

Magnetic Polarization of Light. The phenomena in which a beam of plane polarized light is rotated when passed through a transparent medium placed in a magnetic field. Mathematically,

$$w = \phi H l$$

where w=the angle of rotation, l=the length of the transparent medium, H=the magnetic field strength, and ϕ =Verdet's constant.

Magnetic Quantum Number, m. Determines the orientation of the electron orbital in a magnetic field and has values from +1 to -1, where 1 is the secondary or azimuthal quantum number. It also explains the Zeeman effect of splitting up of spectral lines when the source emitting the radiation was placed in a strong magnetic field. It also determines the component of the angular momentum in the direction of the magnetic field for any spatial orientation of the orbit of the electron and such

components have momentum values of m. $\frac{h}{2\pi}$.

Magnetic Separation. A process by which minerals can be separated into ferromagnetic, magnetic, diamagnetic and non-magnetic fractions.

Magnetic Susceptibility. The extent of magnetization of any substance is a function of the field in which the substance is

placed. The magnetic susceptibility per unit volume (K) is

 $K = \frac{I}{H}$

where I=the intensity of magnetization induced, and H=the

The molar or molecular susceptibility of a substance is denoted by χ_m and is given by $\chi_m = K'(M/\rho)$

where M=gram molecular weight and p=the density determined by Gouy balance but nuclear magnetic resonance method can also be used. For paramagnetic substances, $\frac{1}{\gamma} \propto T$ (absolute temperature).

Magnetic Tape. Made of magnetic particles (e.g., γ-Fe₂O₃) on which information is imprinted via electrical signals converted to magnetic fields, which in turn affect the magnetization of the

Magnetism. A study of magnetic force, fields and their effects on other substances. Magnetic fields are produced by moving charge (current in a coil-electromagnet) on a large scale. Different types of magnetic behaviour result from the type of

- Diamagnetism is due to the orbital motion of electrons of those elements whose atoms have no unpaired electrons.
- Paramagnetism is due to the electron spin and a property of elements having unpaired electrons.
- Ferromagnetism also involves electron spin. A few elements like iron are attracted into a magnetic field very

Magnetite, Fe₃O₄. An iron ore. Used as a flux, a pigment for glasses and as a lining material in furnaces.

Malachite. A green mineral consisting of hydrated copper carbonate, Cu₂(OH)₂CO₃. It is used as an ore and a pigment

Maleamic Acid, C₄H₅NO₃. M.P. 172-173°C. The half amide of

Maleic Acid, cis-butanedioic Acid, C4H4O4.

H.C-COOH H.C-COOH

Colourless crystals soluble in water and alcohol. Prepared by treating maleic anhydride with water. On prolonged heating at

150°C it is converted to fumaric acid (an isomer). Reduced by hydrogen to succinic acid. Oxidised by alkaline $KMnO_4$ to mesotartaric acid. Used in the preparation of (\pm) -make acid.

Maleic Anhydride, $C_4H_2O_3$, \parallel HC-CO O. Colourless crystal, HC-CO

M.P. 53°C. Soluble in acetone and CHCl₃. Manufactured by the oxidation of benzene or furfural by air in the presence of vanadium catalyst. Forms maleic acid with hot water. Used in the manufacture of synthetic resins, as antioxidant of fats and oils during storage and to prevent the development of rancid odours in pastry and milk powders.

Malic Acid, Hydroxy Succinic Acid, C₄H₆O₅. Contains an asymmetric carbon atom, and exists in two optically active and a racemic form. (—)-Malic acid crystallizes in colourless needles, M.P. 100°C. Occurs in many acid fruits (e.g., apples, grapes and goose berries). Prepared from (+) bromosuccinic acid and NaOH.

Malleability. Describing a property in which a body (mainly a metal) can be extended in all possible directions by hammering or rolling.

Malonic Acid, C₃H₄O₄. Colourless crystals CH₂ COOH

Soluble in alcohol and water. Manufactured by heating a solution of sodium cyanoethanoate with NaOH until no more NH₃ evolved. Decomposes above 140°C to form acetic acid. Its esters are used in organic synthesis.

Malonic Ester, Diethyl Malonate, C7H12O4,

COO CH₂CH₃
COO CH₂CH₃

A colourless liquid with a faint aromatic odour, B.P. 199°C. Miscible with organic solvents. Prepared by treating sodium monochloroethanoate with NaCN in alkaline solution at 60°C. The sodium cyanoethanoate is heated with alcohol and H₂SO₄ to form malonic ester which is converted to malonic acid and alcohol by boiling with water or dilute alkalis. It contains a reactive methylene group (CH₂<). The hydrogen atoms in the methylene group are reactive due to their position between

168 Malonitrile

two negative groups. One of these is readily replaced by sodium when treated with alcoholic solution of sodium ethoxide. Used for the synthesis of various organic substances, e.g., fatty acids, dibasic and polybasic acids, αβ-unsaturated acids and cyclic and ring compounds.

- Malonitrile, CH2 (CN)2. Used in the synthesis of vitamin B.
- Cereal grain which has been allowed to germinate and then heated to destroy vitality and dried. The sprouting activates the enzyme system. Malt is prepared from barley, wheat rye, corn, rice, etc. Used in bearing and in food additives.
- Maltase. An enzyme present in the digestive system of animals and in yeast. It hydrolyses the sugar of starch (maltose). It is also present in malt which specifically splits maltose into two mole-
- Maltose, C₁₂H₂₂O₁₁. A sugar found in germinating cereal seeds. A disaccharide composed of two glucose units. It is hydrolysed to glucose. It is produced by degradation of starch and glycogen. It is 4.[a-D-glucopyranosido]. [D-glucopyranose], the a-form is monohydrated, M.P. 102-103°C. Used in brewing, soft drinks and food.
- Mandelic Acid, 1-hydroxy (1-phenyl), Ethanoic Acid, C₈H₈O₃,

Colourless rhombic prisms, (±) M.P. 118°C,

(+ or -) M.P. 133°C. Soluble in alcohol and ether. Prepared by hydrolysing benzaldehyde cyanohydrin. Used in urinary infections.

- Manganates (VII), [MnO₄] (Permanganate). Purple salts, obtained by electrolytic oxidation of [MnO₄]²⁻ Strong oxidizing agents.
- Manganates (VI), [Mr.O₄]²⁻. A salt containing the ion [MnO₄]²⁻. Obtained from MnO₂, fused KOH and KNO₃, air. Deep green
- Manganates (IV), magnesites. Mixed metal oxides having Mn(IV).
- Manganates (III), Mn₃O₄. Mixed metal oxides.
- Manganese, Mn. At. No. 25, At. wt. 54.93. D 7.2. A grey transition metal occurring naturally as oxides, e.g., pyrolusite (MnO₂). An element of group VII with electronic configuration $1s^2$, $2s^2$ p^6 , $3s^2$ p^6 d^5 , $4s^2$. Occurs in nodules on the sea-bed. Prepared by roasting the ore followed by reduction with Al or C and finally by electrolytic reduction. Manganese decomposes cold water and dilute acids to produce hydrogen and reacts with O_2 and N_2 on heating. The oxidation states are +7, +6, +4 and

(the most stable) +2. Manganese (II) salts are pale pink, Mn(II) compounds are brown, Mn(IV) compounds are strong oxidising agents.

- Manganese Chlorides. The only stable chloride is MnCl₂. It crystallizes at the ordinary temperature as the α-tetrahydrate, MnCl₂, 4H₂O. Below -2°C a hexahydrate, and above 58°C a dihydrate are formed. Anhydrous salt may be obtained by heating the hydrates in a steam of hydrogen chloride. Soluble in water.
- Manganese Chlorides Complex. Two series are known. For example, potassium mangnichloride (K₂MnCl₅), (NH₄)₂ MnCl₅, H₂O, Rb₂MnCl₂, etc., and chloromanganites (K₂MnCl₄,Rb₂. MnCl₆ etc.)
- Manganese Oxides. (a) Manganese (II) Oxide, MnO (manganous oxide). A green powder prepared by heating manganese (II) carbonate or oxalate in the absence of air. A basic oxide, insoluble in water, reduced by hydrogen to manganese.
 - (b) Manganese (III) oxide, Mn_2O_3 (manganic oxide, manganese sesquoxide). A black powder prepared by igniting manganese (IV) oxide or a manganese (II) salt in air at 800°C. Reacts with cold acids to form salts. Occurs as braunite (3Mn₂O₃, MnSiO₃) and as the monohydrate (Mn₂O₃ H₂O). Its lattice contains Mn³⁺ and O₂ ions.
- Manganese (IV) Oxide, MnO₂ (manganese dioxide). A black powder prepared by heating MnNO₃. Occurs as hydrated form in pyrolusite. A powerful oxidizing agent, reacts with HCl and H₂SO₄ to produce Cl₂ and O₂ respectively. Alkaline KMnO₄ forms dihydrate (MnO₂, 2H₂O) with MnO₂. Used as a catalyst (in Cl₂ preparation), as a depolariser in dry cells and glass industry.

Manganese (II) Phosphate. Formed by precipitation from aqueous solutions, e.g., Mn₃(PO₄)₂, 7H₂O. Used for the estimation of manganese.

Manganese Sesquoxide. See manganese (III) oxide.

Manganic Salts. Commonly Mn (III) salts.

Manganites. Generally Mn (IV) salts.

Mannans. Polysaccharides consisting of a chain of mannose unit.

Mannich Reaction. The amino ethylation of organic compounds containing active hydrogen atoms. It is carried out by reacting 37% methanal (CH₃OH) with the amine, e.g., (CH₃)₂NH, and the organic compound containing active hydrogen atom (e.g., RH) or atoms.

 $(CH_3)_2NH+CH_2O+RH\rightarrow R.CH_2. N(CH_3)_2+H_2O$

- Manninotriose, C18H32O16 (glucose galactose galactoside). Present
- Mannitol, C₆H₁₄O₆ (alcohol of mannose). Present in fungi and plants. Optically active.
- D-Mannose, C₆H₁₂O₆. A simple sugar found in many polysaccharides. M.P. 199°C. It is an aldohexose, isomeric with glucose.
- Manometer. A device for measuring pressure.
- Marble, CaCO₃. A dense mineral form of calcium carbonate, ob-
- Margarine. A substituted food product for butter, obtained from vegetable oils (polyunsaturated fats).
- Markownikoff's Rule. A rule stating that in the ionic addition of an unsymmetrical reagent (H-G), e.g., (HCl), the hydrogen or positive end of the reagent becomes attached to the carbon atom of the double bond bearing the large number of hydrogen atoms. For example, $(CH_3)_2C=CH_2$ can form two addition products with HX, namely, $(CH_3)_2$ HC. CH_2X or $(CH_3)_2X$ C.CH₃. But according to the rule the product obtained is

The rule has an exception in case of free radical reactions.

Marsh Gas. Methane.

- Marsh's Test for Arsenic. Arsenic containing specimen is converted to volatile AsH₃ (arsine) by nascent hydrogen which in turn, is decomposed to a brown mirror by heating. Estimation of arsenic by comparing with standard mirror. The stain is soluble in sodium hypochlorite, a distinction from Sb which
- Maser. A device working on laser principle but with microwave
- Masking. A technique (analytical) in which addition of a masking agent prevents interfering species from taking part in the
- Mass, m. A measure of the quantity of matter in a object. S.I.
- Mass Action, Law of. A law stated by Guldberg and Waage (1864) which states that the rate at which a substance reacts is proportional to its active mass and the overall rate of the reaction is proportional to the product of the active masses of the

the velocity or rate of forward reaction $v_1 \propto [A][B]$ or $v_1 = k_1[A][B]$.

where ki is a specific reaction rate constant and [A] [B] are the active masses of reactants at equilibrium.

Similarly,

Velocity of the backward reaction $v_b = k_b$ [C] [D], where k_b is also a reaction rate constant and [C] [D] are active masses of products at equilibrium.

At equilibrium,

$$v_f = v_b = k_f[A] [B] = k_b[C] [D]$$
or
$$\frac{k_f}{k_b} = \frac{[C] [D]}{[A] [B]} = K$$

where K is the equilibrium constant.

In actual practice concentration terms in brackets represent active concentrations instead of active masses, because active mass=activity coefficient×active concentration.

But for dilute solutions or for gaseous reactions, activity co-efficient is taken as unity. Hence, [A] [B] or [C] [D] are the molar concentrations of reactants and products.

Mass Defect. See packing fraction. Massicot, PbO. See lead oxide.

Mass Energy Equation.

E=mc2

where E=the total energy (rest-mass energy+ kinetic energy+ potential energy), m=mass and c=the velocity of light. The equation is a basis of Einstein's Theory of Relativity. Energy and mass are interconvertible. Conversion of rest-mass into energy (kinetic) is the source of power in radio-active species.

Mass Number. The number of protons+number of neutrons=
The mass number is the nucleus.

Mass Spectrography. A refined modification of the parabola method of positive ray analysis. In the mass spectrometer the relative abundance of isotopes or the other ions is determined by measuring the positive ion currents received by an electrometer with several controlled magnetic fields and accelerating potentials.

Mass Spectrometer. See mass spectrograph.

Matches. A device for producing flame. Matches consist of potassium chlorate and red phosphorous with an active material such as glue matrix and a glass powder (the match stick).

172 Matlockite

Matlockite, PbFC1.

Matrix. A continuous solid phase in which particles of a different solid phase are embedded.

Matte. An indefinite mixture of iron and copper sulphides obtained in smelting copper ores.

Maxwell, Mx. A unit of magnetic flux in C.G.S. units and equal

MCD. Magnetic circular dichroism.

MCPA, 2-methyl-4-chlorophenoxyacetic Acid, Methoxone, C₉H₉ClO₃. M.P. 118°C. A high brown solid. selective weed-killer. Used as a

MCPB, 4-(4-chloro-2-methylphenoxy) butyric Acid, C11H13ClO3. A compound which converts into a power herbicide in plant cells. Used as a selective weed-killer.

Md. Mendelevium.

MDI. Methylene di isocynates.

Me. Methyl, CH₃-.

Mean Free Path, (λ). The average distance travelled by a molecule between two successive collisions of a molecule.

Mechanism, of a Reaction. The mode by which a chemical reaction occurs. Mechanism is expressed by suitable chemical equations. For example, one step reaction, multi-step reaction,

Medicinal Paraffin Oil. Liquid paraffin.

Mega, (M). A prefix denoting 106, e.g., 1 megahertz (MHz)=

MEK. Methyl ethyl ketone.

Melamine, Cyanuramide, C₃H₆N₆. M.P. 354°C. An important material used in plastic industry. Condensed with formaldehyde and other substances it produces thermosetting resins which are

Melanin. A dark brown pigment of hair, eyes and skin. A colloidal amorphous substance which is a polymer obtained by the enzyme

Melibiose, C₁₂H₂₂O₁₁. A hydrolysis product of raffinose.

Melissic Acid. Triacontanoic Acid, C₃₀H₆₀O₂, CH₃ [CH₂]₂₈ COOH.

Melissyl Alcohol Myricyl, Alcohol, C₃₀H₆₂O, CH₅ [CH₂]₂₈. CH₂OH. M.P. 87°C. Present in bees' wax.

- Mellitic Acid, C₁₂H₆O₁₂. M.P. 286-288°C. Occurs as Al-salt in some lignite beds. Obtained by charcoal and conc. HNO₃.
- Melting Point. The temperature at which a solid melts and is in equilibrium with its liquid at a standard pressure.
- Membrane. A thin pliable sheet of tissues or other materials acting as a boundary. Two types of membranes, e.g., natural such as in cells and skins and synthetic, e.g., cellulose derivatives or rubber.
- Membrane Cell. An electrolytic cell used for the production of NaOH from brine. The anode and cathode of the cell are separated by a membrane. For example, Castner-Kellner cell and mercury cell.
- Membrane Hydrolysis. A process in which a colloidal electrolyt; separated from water by a membrane (dialyser) becomes either acid or alkali inside the dialyser.
- Mendelevium, Md. At. No. 101. Two isotopes ²⁵⁶Md and ²⁵⁸Md. Obtained by bombardment of Es with ⁴He is a cyclotron. Forms no solid compounds.
- Mendius Reaction. The reduction of the cyanide group to a primary amine group using Na/alcohol:

$RCN+2H_2\rightarrow RCH_2NH_2$

A method of increasing the chain length of compounds in ascending a homologous series.

Menthol
$$C_{10}H_{20}O$$
, CH_3

An optically active monocyclic terpene alcohol. A constituent

 CH_3
 CH_3

of peppermint oils. (—)-Menthol is a solid which crystallises in four different forms (M.P. 43°C). Obtained by reduction of menthone, isomenthone or piperitone. Used externally as an analgesic in rheumatism and sinusitis.

Menthone, C₁₀H₁₈O. An optically active mono cyclic ketone. Contains two asymmetric carbon atoms and forms 4 optically active and 2 racemic modifications. (—)-menthone is prepared by oxidizing menthol. Present in some oils.

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- Mercaptals, RR'C (SR")2. Oily liquids acetals in which the oxygen atoms have been replaced by sulphur atoms. Obtained from mercaptans and carbonyl compounds in the presence of zinc chloride or HCl. Used for protecting the carbonyl group.
- Mercaptans, Thiols. Organic compounds containing an -SH group directly linked to a carbon atom. Also the alcohols in which the oxygen atom has been replaced by sulphur. Liquids with unpleasant odour. Present in crude petroleum. Prepared from alkyl or oryl halides and KHS. Used to control a polymer chain, as odorants, as rubber vulcanizer, etc.
- Mercapturic Acids. Derivatives of N-acetylcysteine excreted in the
- Mercurammine Compounds. A group of compounds obtained when ammonia gas reacts with several chemical species. For example, compound HgCl₂, 2NH₃ (ammonia gas+mercuric chloride), infusible white precipitate of NH₂. HgCl (ammonia+mercuric chloride). A brown precipitate of NH₂. Hg. O. Hg. I (ammonia+Nesslers' reagent), a yellow powder, Millon base NHg2. OH, 2H₂O (mercuric oxide on warming with aqueous ammonia). On drying at 125°C in ammonia gas, the dark brown explosive compound NHg2OH is formed.
- Mercuration. The process by which Hg (II) derivatives of aromatic compounds are formed by direct substitution, e.g., mercury derivatives of benzene (C₆H₅ HgO₂C CH₃).

Mercuric. Mercury (II) compounds.

Mercurous. Mercury (I) compounds.

- Mercury, Hg. At. No. 80, At. wt. 200:59, M.P. 38:87, B.P. 356:58°C, D 13.6. A transition metal occurring naturally as cinnabar, mercuric sulphide. It is a silvery white liquid metal. Its electronic configuration is $5d^{10}$, $6s^2$. Vapour density corresponds to the monoatomic formula Hg. Attacked by hot conc. H₂SO₄ and dilute HNO₃. The normal oxidation state is +2. Exists as (Hg-Hg)2+ ion and as other polymercury cation such
- Mercury Amalgams. Mercury dissolves many metals to form amalgam, e.g., NaHg, NaHg, etc. Copper and cadmium amalgams are used as dental stoppings and cadmium amalgam is used in Weston cadmium cell.
- Mercury Cell. A voltaic or electrolytic cell in which one or both of the electrodes consists of mercury or an amalgam. For example, Daniell cell and the Weston cadmium cell have amalgam electrodes whereas, flowing mercury electrodes are used in electrolytic cells (e.g., in the manufacture of Cl₂).

- Mercury (I) Chloride, Hg₂Cl₂ (mercurous chloride, calomel). A white precipitate (dilute HCl+mercury (I) salt). It is sparingly soluble in water and is blackened by both ammonia gas and by alkalis. Used as a purgative.
- Mercury (II) Chloride, HgCl₂ (mercuric chloride). A colourless crystalline compound (direct combination of mercury and cold dry chlorine). It is soluble in water and dissolves in conc. HCl due to the formation of complex ions, HgCl₄²⁻ and HgCl₃⁻. Sublimes on heating to form a white translucent mass. Used as an insecticide and as a purgative and in the calomel electrode.
- Mercury (I) Iodide, Hg₂I₂. Pale green, precipitated from a solution of a Hg₂²⁺ salt by I⁻.
- Mercury (II) lodide, HgI₂. Scarlet up to 126°C or yellow substance. Prepared from HgCl₂ and KI solution or from Hg and I₂. Forms complexes with I₂, e.g., Nessler's reagent.
- Mercury (I) Nitrate, Hg₂ (NO₃)₂. Crystallizes as a monohydrate from the solution (Hg+cold dilute HNO₃).
- Mercury (II) Nitrate, Hg (NO₃)₂. Formed by dissolving mercury in excess concentrated HNO₃, on crystallization, deliquescent colourless crystals, 2Hg (NO₃)₂, H₂O obtained.
- Mercury, Organic Derivatives. Organe-mercury (II) derivatives such as C_6H_5 HgO₂ CCH₃ [C_6H_6 +Hg (O₂CCH₃)₂]. No mercury (I) derivative is known. Used to prepare other organometallic derivatives.
- Mercury (II) Oxide, HgO (mercuric oxide). A yellow-orange poisonous powder formed by the addition of Hg²⁺ salt to NaOH or by heating mercury to 300°C (a red solid). On strong heating it forms mercury and oxygen. Mercury (I) oxide does not seem to exist.
- Mercury (II) Sulphate, Hg₂SO₄. Colourless solid (Hg+excess hot H₂SO₄). Easily hydrolysed to Hg₂OSO₄ (turpeth mineral).
- Mercury (I) Sulphate, Hg₂SO₄. Obtained by excess Hg+H₂SO₄.
- Mercury (I) Sulphide, Hg₂S (mercurous sulphide). Some doubt about the existence of mercurous sulphide.
- Mercury (II) Sulphide, HgS (mercuric sulphide, vermalia). Occurs in nature as the mineral cinnabar (a red solid) and metacinnabar (black solid). Prepared as a black precipitate by H₂S+a soluble mercury salt or by grinding Hg+S. Used as a pigment.
- Mesaconic Acid, Methyl Fumaric Acid, C5H6O4,
 - H₃C-C-COOH

 M.P. 204°C. Colourless crystals.

 HOOC-C-H

Mescalin, C₁₁H₁₇NO₃. M.P. 35°C. An alkaloid.

Colourless liquid, B.P. 165°C. Occurs in crude petroleum. Prepared from propane plus concentrated H2SO4.

Meso ionic. A five-or-six membered neterocyclic compound which cannot be represented satisfactorily either by covalent or polar structure, and possess a sextet of electrons in association with all the atoms constituting the ring.

Meson, Mesotron. A subatomic particle (responsible for exchanges between neutron and proton) with either a positive, negative or zero charge. Found in cosmic rays. High energy nuclear collisions also produce mesons. Mass ca 200 times that of an electron and mean life time ca 10⁻⁷ seconds.

Mesothorium. Ra and He isotopes formed in natural decay series.

Mesoxalic Acid, Ketomalonic Acid, C3H2O5

Occurs as dihydroxy malonic acid, M.P. 121°C. Prepared from HNO₃ plus malonic acid. Gives some reactions of both a ketone and an acid, e.g., forms phenylhydrazones and esters with phenyl hydrazine and alcohols respectively.

Mesyl. The methane sulphonyl group.

Meta, (m-). A nomenclature (prefix) used only for disubstituted derivatives of benzene, e.g., 1,3-dimethyl benzene is m-xylene.

$$\begin{array}{c} \text{CH}_3 \\ \text{1} \\ \text{6} \\ \text{O} \\ \text{3} \\ \text{5} \\ \text{4} \\ \text{CH}_3 \\ \text{m-Xylene} \\ \text{or} \\ \text{m-Xylene} \\ \end{array} \begin{array}{c} \text{CH}_3 \\ \text{6} \\ \text{O} \\ \text{2} \\ \text{5} \\ \text{3} \\ \text{4} \\ \text{Xylene} \\ \end{array}$$

The same prefix is found in metaphosphoric acid and metabisulphites, but in such cases it is written in full.

Meta-position is always the third position of carbon with respect to carbon number one in benzene ring whereas first and fifth positions of carbon are denoted by o-(ortho) and p-(para) Metallocene 177

Metabolism. The chemical reactions which take place in cells. The molecules taking part in metabolic reactions are called metabolites. Metabolic reactions involve the breaking down of molecules to provide energy (catabolism), e.g., breakdown of sugar to CO₂ and water and release of large amounts of energy. In anabolism, a synthetic reaction takes place, e.g., the formation of fats or proteins.

- Metaboric Acid, HBO₂. An intermediate product during the loss of water from boric acid.
- Metal. Elements which have a characteristic lustre, good conductivity (for heat and electricity), and undergo chemical reactions as positive ions or cations, e.g, Cu, Ag, Na, Hg, etc.
- Metalation. A process of removing a relatively acidic proton from a compound and replacing by a metal atom such as Na or K.
- Metal Carbonyls. Co-ordination compounds of CO and a metal (e.g., transition metals) in which the carbon is usually bonded to the metal. The general formula may be written as:

M(CO)n, where n=1,2,3, etc.

Most of the transition metals form stable carbonyls. Simple stable carbonyls, except $V(CO)_6$, have an electronic configuration corresponding to the next noble gas.

- Metal Cluster Compounds. Compounds made up of clusters of metal atoms linked together by covalent or co-ordinate bond.
- Metaldehyde, $(C_2H_4O)_n$ (n=4 or 6). A solid, M.P. 112-115°C. Used as a fuel and slug control.
- Metallic Bond. A bond formed between atoms of a metallic element in its zero oxidation state in an arrangement of similar atoms.
- Metallic Conduction. The conduction in metals is due to the movement of electrons without decomposition. Each atom in the metal contributes only one or two electrons to the energy state termed as 'the conduction band' and these are the electrons which are free to move.
- Metallic Soaps. Water insoluble, alkaline earth, heavy metal, Li salts of long chain carboxylic acids. Napalm is an Al soap [Al (OH) R₂] used in greases, lubricating oils, rubber etc.
- Metallizable Dyes, Mordant Dyes. Dyestuffs with a metal atom added in situ after application to fibre.
- **Metallocene.** A sandwich compound in which a metal atom is coordinated to two cyclopentadienyl ions, e.g., $[Fe(C_5H_5)_2]$ of $[(C_5H_5)_2Co)$. The bis- $(h^5$ -cyclopentadienylmetal) compounds.

- Metallochromic Indicator. A compound that can form a complex with a total different colour from that of the free indicator, e.g., eriochrome black T used with Mg2+.
- Metalloids. Materials which are intermediate in properties between metals and non-metals, e.g., germanium, arsenic and
- Metallurgy. The study of the extraction, working and use of
- Metal-metal Bonds. Bonds (covalent or coordinate), between
- Metal Passivators. Poisoning or passivation of a catalyst in a reaction by small traces of metals (passivators).
- Metal Surface Treatment. The removal of oxides (chemically). dirt and grease (by organic solvents) followed by pickling in acid
- Metamerism. A change in the observed colour.
- Metaphosphates. Phosphates having cyclic or chain linked PO43-
- Methacrylic Acid, 2-Methylpropenoic Acid, C₄H₆O₂. Colourless solid. M.P. 15-16°C. B.P. 160.5°C. Used in the preparation of synthe-
- Methadone C₂₁H₂₇NO. A powerful analgesic ketone.
- Methanal, HCHO (formaldehyde). A colourless gaseous aldehyde.

 Manufactured by the oxidation of methanol (500°C and Ag-

$2CH_3OH + O_2 \longrightarrow 2HCHO + 2H_2O$.

A solution of methanal (40%) in water is known as formalin which is extensively used as a preservative for biological specimens. Forms polymethanal when an aqueous solution of methanal is evaporated (-O-CH₂-O-CH₂-O-CH₂-).

- Methane, (CH₄). A gaseous alkane (saturated) also known as 'Marsh Gas'. M.P. -184°C, B.P. -164°C. Occurs in coal mines where its mixture with oxygen is termed as "Firedamp'. Manufactured from CO (1 volume) and H₂ (3 volume) passing over heat Ni catalyst at 230-250°C. Combines explosively with Cl₂ at room, temperatures; at low pressures methyl chloride is formed. Used in the manufacture of H₂, NH₃, CO, carbon black
- Methanesulphonic Acid, CH₃SO₂OH. M.P. 20°C, B.P. 167°/10 mm. Soluble in water but insoluble in hydrocarbons. Obtained by oxidation of dimethyl sulphide. Used as a catalyst in esterification, dehydration, polymerization and alkylation reactions.

Methanoates, Formates. Esters and salts of methanoic acid.

Methanoic Acid, HCOOH (formic acid). A liquid carboxylic acid (colourless) fuming slightly. M.P. 8'4°C, B.P. 100'5°C. Obtained by action of sulphuric on sodium methanoate (NaOOCH). A strong reducing agent that occurs in ants. Used in textile dyeing and finishing, in leather tanning and a solvent for organic and inorganic compounds.

Methanol, Methyl Alcohol, Wood Spirit, Wood Naptha CH2OH. colourless liquid alcohol soluble in water. B.P. 64'5°C. Poisonous and causes blindness. Occurs as ester in various plants (e.g., winter green) and manufactured by the catalytic oxidation of methane from natural gas.

$$CH_4 \xrightarrow{[O]} CH_3OH$$

Oxidized in air to methanal and reacts with sulphuric acid to form methyl hydrogen sulphate (CH₃HSO₄), dimethylsulphate [(CH₃)₂SO₄] and dimethylether [(CH₃)₂O). Reacts with Na to from sodium methoxide. Solvent for inorganic salts and organic compounds. Used in the manufacture of methanoic acid and MTBE (5%).

Methenamine. Hexamine.

Methionine, 2-Amino-4-(methylthio)butanoic Acid, C₅H₁₁NO₂S,

CH₃SCH₂CH₂CH(NH₂)COOH. M.P. 283°C. A naturally occurring sulphur containing aminoacid (-).

Methoxy Group. The group CH3O-.

Methylal, Methylformal, Dimethoxymethane, C3H8O2.

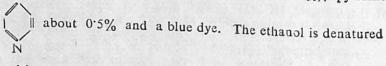
Colourless liquid, B.P. 42°C. Occurs in commercial formalin. Used as a solvent and a substitute for methanol in many reactions.

Methyl Alcohol. See methanol.

Methyl Amines. Compounds of ammonia (NH₃) in which 1, 2, or 3 of the hydrogen atoms have been replaced by methyl groups of the nydrogen atoms in the start groups $(-CH_3)$, e.g., monomethylamine, methylamine (CH_3NH_2) , dimethyl amine $[(CH_3)_2NH]$ and trimethylamine $[(C_3H)_3N]$. Manufactured by the vapour phase reaction of methanol and ammonia at 350-450°C under pressure over an Al catalyst. The product obtained is a mixture of all three compounds.

Form alkaline solutions with water but the alkalinity decreases with the increase in number of -CH₃ group.

Methylated Spirit. Ethanol containing methanol (9.5%), pyridine



and is used as a fuel and a solvent.

Methylation. A reaction involving the introduction of a methyl group (-CH₃), e.g., Friedel-Crafts reactions with halomethanes.

Methyl Bromide, CH₃Br. Bromomethane.

Methyl Cellosolve. Ethylene glycol monomethyl ether.

Methyl Cyclohexane, C₇H₁₄, C₆H₁₁CH₃. Colourless liquid, B.P. 101°C. Used as a solvent.

Methylcyclohexanol, Hexahydrocresol, C₇H₁₄O. A colourless viscous liquid (a mixture of isomers) with a boiling range 165-180°C. (Cresol +H₂ in the presence of Ni). Used as a solvent for fats, oils, gums, etc.

Methylcyclohexanone, C₇H₁₂O. A colourless liquid (mixture of a heated catalyst). Used as a solvent for resins and gems.

Methylene, (=CH₂). Obtained in keten free state by the thermal or photochemical dissociation of keten or diazomethane. Very

Methylene Blue (Basic Blue). A dyestuff prepared by boiling indamine with dilute acid or a solution of zinc chloride. Used as a staining matter, in dyeing fast fibres and a mild antiseptic.

Methylene Diisocyanate, MDI, CH₂(NCO)₂. An intermediate used in the manufacture of urethane foams.

Methyl Ethanoate (methyl acetate), CH₃COOCH₃. A colourless liquid ester having a fragrant odour, used as a solvent.

Methylene ditannin, $CH_3(C_{14}H_9O_9)_2$. A condensation product of HCHO and tannic acid. Used as a dusting powder or an ointment for eczema.

Methyl Ethyl Ketone, C₄H₈O, CH₃COCH₂CH₃. B.P. 78.6°C. A colourless liquid, soluble in water to 24% by weight, dation of sec butyl alcohol with air in presence of heated Cu catalyst. Used as a solvent.

- Methyl Fluorosulphate, FS(O)₂OCH₃. B.P. 94°C, A colourless liquid and a powerful methylating agent (e.g., for amides and nitriles).
- Methyl Glyoxal, Pyruvic Aldebyde, C₃H₄O₂, CH₃COCHO. Yellow liquid, B.P. 72°C (oxidation of propanone with SeO₂).
- Methyl Icdide, CH₃I. (Iodomethane). Colourless liquid, B.P. 43°C. Prepared from methanol plus iodine with red phosphorous (heating). Used for preparing Grignards reagent (methyl) and and as a methylating agent.
- Methyl Isobutyl Ketone, MIBK, Hexone, ketone used as solvent for many polymers. Prepared by selective hydrogenation of mesityl oxide.
- Methylnaphthalenes. Occurs in crude oil and reformed petroleum. Naphthalene carboxylic acids are used as dyestuff intermediates.
- N-methyl-N-nitro-N-nitrosoguahidine, C₂H₅N₅O₃. M.P. 118°C. A potent mutagen for organisms.
- Methyl Orange, 4- dimethylamine-4'-azobenzene, Sodium Sulphonate.

Orange crystals prepared by the action of dimethylaniline on diazotised sodium sulphanilate. Used as an indicator in 0.01% aqueous solution [PH range 3.1 (red) to 4.4 (yellow)].

- N methyl-2 pyrrolidone. Biodegradable solvent used mainly for resins.
- Methyl Red. O-Carboxybenzeneazodimethylaniline, $C_{15}H_{15}O_2N_3$. M.P. $181^{\circ}C-182^{\circ}C$. An indicator,

prepared by reacting dimethylaniline and diazotized anthranilic acid. Used in 0.02% solution in 60% alcohol [pH range 4.4 (red) to 6.0 (yellow)].

- Methyl Phenols, Cresols, HO.C₆H₄CH₃. Compounds containing both methyl and hydroxyl groups attached to the benzene ring. Obtained from coal tar (a mixture of isomers with the methyl group in the 2-, 3-, and 4-positions respectively).
- Methyl Salicylate, C₈H₈O₃. Colourless liquid, M.P. 8·6°C, B.P. 233°C. Occurs in pure state in essential oils of winter green and sweet birch. Readily absorbed by skin and is prepared by the estesweet birch. Readily absorbed

rification of salicylic acid. Used in perfumery and as a flavouring

Methyl Violet (Basic Violet I). A violet dye obtained by the oxidation of dimethylaniline with CuI₂ and a mixture of the hydrochlorides of tetra-, penta-, and hexa-methyl-p-rosanilines.

Used for dyeing jute, for colouring methylated spirits, as a bacteriological strain and also as an indicator.

Metol. Aminophenols.

Mevalonic Acid, C₆H₁₂O₄. D Mevalonic acid, the 3(R) form of 3, 5dihydroxy-3-methylpentanoic acid was isolated as the δ-lactone from distiller's dried solubles. It is an intermediae in the biosynthesis of the terpenoides and steroids (polyisoprenoids). Mg. Magnesium.

MIBK. Methyl isobutyl ketone.

Several aluminosilicates with layers of linked (Si, Al)O4 tetrahedra which resemble clays and tale. For example, phlogopite (KMg₃(OH)₂Si₃AlO₁₀), and (K₂Mg₆Al₂Si₆O₂₀F) fluorophlogopite (used in electricals). It splits into thin insulating sheets.

Micelle. Describing aggregated colloids, e.g., soap and dyes.

Michaelis Constant. 'The substrate concentration at which the rate of reaction is half its maximum rate' is called Michaelis

Michael Reaction. A reaction used to define the conjugate addition of nucleophiles across double bond conjugated with electronwithdrawing groups such as carbonyl (>C=0) in CH₈CH=

Michler's Hydrol, bis (p-dimethylaminphenyl) Carbinol, C₁₇H₂₂N₂O. M.P. 96°C. Colourless crystals. Used in the preparation of

Michler's Ketone. C₁₇H₂₀N₂O. M.P. 172°C, (Phosgene+Dimethyl-

Microbalance. A balance used for weighing in order of 10⁻⁶ gram

Microcosmic Salt, NaNH₄HPO₄, 4H₂O. See sodium ammonium

Microcrystalline Wax. See petroleum wax.

Micron, (μ) . 1 micron=10⁻⁴ cm.

Microstate. Describing all the possible arrangements in which the total energy of particles of a system may be distributed among the particles. Each of these possible arrangements is termed as microstate.

Microwave. A form of electromagnetic waves with wavelength (λ) in the range 10⁻² to 1 mm or frequency 10¹⁰ to 10¹² Hz. Microwaves are produced by various electronic devices including the klystron; they are often carried over short distances in tubes of rectangular sector called wave guides. Physical effects—heating used for food and polymerization can be induced by microwaves.

Microwave Spectroscopy. Spectra are obtained by passing microwave beam (klystron method) through the gaseous sample; the transmitted beam being detected by a crystal receiver. The spectra correspond to transition between levels very close in energy and provides information on the rotational energy levels of some molecules. Used for the identification of molecules in the interstellar medium.

Milk of Lime. A suspension of calcium hydroxide in water.

Millimicron, $m\mu$. I millimicron = 10^{-7} cm.

Mineral Colours. Inorganic pigments used for colouring paints, plastics and papers, e.g., prussian blue, iron buff, etc.

Mineral Solvents. White spirits.

Minium. Pb3O4.

Miscibility. A term used to describe the extent of mixing, e.g. completely miscible (C₂H₅OH+water), partially miscible (phenol+water), and immiscible (C₆H₆+water).

Mispickel Arsenopyrite, FeAsS. An important are of As.

Mixed Crystals. (Solid solutions). Crystals consist of a mixture of the two substances from the solution of which they have been crystallized. The substances present in a solution crystallise in similar form.

Mixed Indicator. A mixture of two or more indicators used to restrict the pH range or sharpen the end point.

Mixed Metal Oxides. Compounds derived from oxides and which contain two or more metals in an arbitrary ratio obtained by heating mixtures of oxysalts.

Mixture. Two or more substances when mixed form a mixture in which there is no chemical bond. Homogeneous mixture

has only one phase, e.g., mixtures of gases or solutions. Heterogeneous mixture can have two or more than two phases, e.g., some alloys or a mixture of sugar and sand. Mixture differs from a chemical compound in the following respects:

- The chemical properties of the mixture are same as those
- It can be separated by physical means, e.g., distillation or 2.
- The proportions of the component in a mixture can vary, 3.
- M.K.S. System. A system of units based on the meter, the kilogram
- mm Hg (millimeter of mercury). A former unit of pressure and is equal to 133.3224 Pascal and is almost identical to torr.

Mn. Manganese.

Mo. Molybdenum.

Mobility, Ionic. The absolute velocity of migration of ions, the ionic mobility (u), which are the mean velocities of the ions in cms per second under a potential gradient of 1 volt per second.

$$u_{+} = \frac{\lambda_{+}}{F}$$
 and $u_{-} = \frac{\lambda_{-}}{F}$

where λ_{+} and λ_{-} are the equivalent conductivities of cations and anions at infinite dilution ($\lambda_{\infty} = \lambda_e + \lambda_a$, Kohlrausch's Law).

Mehr's Method. A method for the volumetric estimation of Cl with Ag+ in the presence of CrO42 indicator (a red ppt. at the end point).

Mohr's Salt, (NH₄)₂ SO₄, FeSO₄. 6H₂O. Iron ammonium sulphate.

Molality (m). Mode of expressing the concentration of a solution in terms of number of moles of solute dissolved in 1000 grams

Molar Conductivity. The conductivity of a solution of an electrolyte containing 1 mole of the electrolyte and is given by

where μ =the molar conductivity, K=the specific conductivity and V=volume in millilitres containing 1 mole of the electrolyte.

Molar Heat. The amount of heat required to raise the temperature of 1 mole of a substance by 1°C, at either constant pressure of Molecule 185

Molarity (M). The concentration of a solution expressed as the number of moles of solute present in 1 litre of solution. A 1M solution of NaOH contains 40 g. of the solute in 1 litre of solution.

- Mole (mol). The amount of substance that contains one gram formula weight of the substance. One mole of a substance contains molecules equal to Avogadro's number, i.e.: 6.023×1023. Also known as one gram molecule.
- Molecular Beam. A beam of molecules in which all the molecules possess velocities as well as energies within a very narrow range. Used in the measurements of molecular velocity distributions and molecular collision cross-sections.
- Molecular Crystals. Crystals in which molecules occupy lattice points, e.g., solid carbon dioxide (dry ice) or iodine.
- Molecular Diameters. For the purpose of chemical kinetics the molecules are supposed to be spherical. The diameters of some molecules are as follows:
 - $H_2~2.38\times 10^{-8}~cm$; $O_2~3.19\times 10^{-8}~cm$; $C_6H_6~6.6\times 10^{-8}~cm$.
- Molecular Distillation. A distillation carried out at a low pressure (1.3 N/m²). Used in the separation and purification of vitamins and other natural products. Also used to separate the isotopes of mercury.
- Molecular Orbitals. Orbitals which belong to all atoms forming the molecule. Only outer electrons are considered for molecular orbitals. Molecular orbitals are classified as:

bonding orbitals-the electron tries to hold the molecules together.

anti-bonding orbitals—the electron causes disruption of molecule. non-bonding orbitals—the electron has no bonding effect on molecule.

- Molecular Refractivity. See specific refractivity.
- Moelcular Sieve. Molecules containing channels, e.g., zeolites used for adsorption of water or gases. Sieves may contain pores of different sizes. Molecular-sieve chromatography is widely used in chemistry and biochemistry.
- Molecular Spectrum. The absorption or emission spectrum which is a characteristic of a molecule. Molecular spectra are generally band spectra.
- Molecule. The smallest particle of matter which can exist in free state, e.g., the ionic substance KCl is considered to be a molecule.

Molecular, Mass of. A molecule (m) is

Gram molecular weight Avogadro's number .

So the molecular mass of a oxygen molecule is $32/6.02 \times 10^{23}$

Mole Fraction. A mode of expressing the concentration of a mixture. If a mixture (say solution) has two components A and B, then XA the mole fraction of A in the mixture is given by

$$X_A = \frac{n_A}{n_A + n_B}$$

and the mole fraction of B, $X_B = \frac{n_B}{n_A + n_B}$

where nA and nB are the numbers of moles of A and B in the

Molisch's Test. A general test for carbohydrates. The carbohydrate solution+alcoholic a naphthol+concentrated sulphuric acid from the sides of the tube - formation of a deep violet ring

Furfural test (distinction test for glucose and fructose). Mixture of the sugars + a-naphthol + concentrated HCl followed by boiling-appearance of violet colour immediately in case of fructose and slow appearance of colour for glucose.

Molybdates. Similar to tungstates, e.g., [Mo₇O₂₄]⁶⁻. Used as-

Molybdenite, MoS2. An ore of Mo. Used as a lubricant.

Molybdenum, Mo. At. No 42, At. Wt. 95.94, M P. 261.7°C. B.P. 4612°C. A transition element of Group VI with electronic configuration 4d⁵ 5s¹. Chiefore molybdenite (MoS₂) which is roasted to MoO₃, converted to ammonium molybdate, purified, heated to MoO₃ and reduced to Mo with hydrogen. A or Na₂O₂. Used in steels and its alloys used for cutting at high temperature. Also used in pigments and as a solid lubricant. The compound ammonium molybdate, dissolved in HNO3 is

Molybdenum shows oxidation states from +6 to -2. Forms

Molybdenum Halides. The important halides are: MoF, MoF, Mo₂F₆, Mo₂F₉, MoCl₆ etc. Complex chlorides are also formed e.g. MoCl₆. Bromide and lodides are similar to Monosilane 187

Molybderum Oxides. Important oxides are: MoO₃, violet Mo₂O₅, brown-violet MoO₂. A mild reduction of hydrated MoO₃. forms a blue hydrous oxide molybdenum blue.

- Molybdenum Steel. 0.2 to 0.3% of Mo added to steel, produces grain refining in its alloys. Molybdenum steel forged readily. Also used as a constituent of stainless and semi-stainless steels.
- Molybdenum Sulphides. MoS₂ show a layer structure and used as a solid lubricant MoS₃ and Mo₂S₃, are other sulphides known.
- Molybdic Acids. A group of acids from which various molybdates are derived. For example, H₂MoO₄ (from MoO₃). Ammonium molybdate [(NH₄)₃ PO₄ Mo₇O₄, 4H₂O] is used for testing phosphates.
- Monazite. A naturally occurring mixed phosphate (Ce, La, Nd, Pr) PO₄ which contains some thorium silicate (ThO₂-1-18%). An important ore of Ce, Th and La.
- Mond Process. A process used to extract nickel from nickel ores.

 See nickel.
- Monel Metal. A nickel alloy containing about 60.70% Ni, 25-35%. Cu, 1-4% Fe, 0.2% Mn and a little Si and C.
- Monobasic Acid. An acid containing only one replaceable hydrogen atom and forms only one series of salts, e.g., HCl and HNO₃, etc.
- Monochromatic Radiation. Radiation of a single wavelength.
- Monochromator. A device used for selecting a single wave length (e.g., prism, grating or a single crystal) from a broad range of radiation or particle. Used in X-ray diffractions and spectrophotometers.
- Monoclinic System. The crystal system characterised by one 2-fold axis of symmetry and/or one plane of symmetry, e.g., Gypsum. A unit cell is comprised of three unequal axis one perpendicular to the other two which are obliquely inclined.
- Monodentate Ligand. A ligand that contains only one co-ordinately active lone pair of electrons which interact through a single atom, e.g., NH₃.
- Monomer. A single molecule, e.g., (CH≡CH) molecule, which can form dimer, trimer or polymer.
- Monosaccharides. Sugars represented by formula $C_nH_{2n}O_n$ (n=5 or 6), e.g., glucose or fructose $(C_6H_{12}O_6)$.
- Monosilane. The silicon hydride SiH₄, a gas, B.P.-111'9°C.

- Monotropy. A form of allotropy in which only one allotropic medification is stable while other forms are metastable and tend to change to the stable modification, e.g., phosphorous violet is stable and its yellow variety is metastable.
- Monovinylacetylene, But-1 ene-3-yne, CH₂=CH−C≡CH. A colour-less gas, B.P. 5°C. Manufactured by the controlled low temperature polymerization of acetylene in presence of aqueous cuprous and ammonium chloride.
- Montanic Acid, Octacosanoic Acid, C₂₇H₅₅COOH. A straight chain saturated fatty acid, M.P. 89°C and occurs in natural wax.
- Montmorillonite. A clay mineral containing aluminosilicate and that has two varieties, e.g., one smells in water to form gels and the other has ren arkable adsorptive power.
- Mordant. An inorganic compound used to dye cloth. The mordant (e.g., aluminium hydroxide or chromium salts) is precipitated in the fibre of the cloth and the dye then gets absorbed in the particles.
- Morphine, C₁₇H₁₉NO₃. Colourless prismatic crystals with one molecule of water, M.P. 254°C (anhydrous). An alkaloid found in opium. Forms codeine and diamorphine or heroin (diacetyl its salts, which are more soluble in water than the base, form a stereoisomer is biologically active. They seem to effect the brain by activating neuronal mechanism.
- Morpholine, C₄H₉ON. A colourless liquid, strong ammoniacal odour B,P. 129°C. Moderately strong base, forms soaps with pholine carbamates. Manufactured by heating ββ'-dichlorodiethylether with ammonia. Used as a solvent and corrosion inhibitor in boilers.

Mosaic Gold. SnS2.

Mossbauer Effect. A phenomena involving the resonance fluorescence by 7-radiation of an atomic nucleus, returning from an excited state to the ground state. Used in the study of Fe, Sn and Sb compounds.

Moseley's Law. Lines in the X-ray spectra of an element have frequencies that depend on the atomic number of the element.

Mathematically,

where v is the frequency radiation, and q and b are the characteristic constants of the element.

Mother Liquor. A solution left after crystallization.

Motor Spirit. A fuel with an approximate boiling range of 310-450 K used for internal combustion engines.

Moulding. A process of giving shapes to objects in metals, plastics, etc.

MSG. Sodium glutamate.

mu-, µ. Designation of a bridging species.

Mucic Acid, 2,3,4,5-tetrahydroxy Hexanedioic Acid, C₈H₁₀O₈.

Colourless solid, M.P. 206°C Obtained by the oxidation of lactose or the galactans from wood with nitric acid. Forms furoic acid and pyrrole. Used to manufacture pyrrole.

Mucilages. Polysaccharides which smell in water (a mixture of galacturnoic acid, xylose and arbinose residues).

Mucopeptides. Insoluble polymers of N-acetylglucosamine and N-acetylmuramic acid or structural components of the cell walls of gram positive and many gram-negative bacteria.

Mussie Furnace. A type of furnace in which charge does not come into contact with the hot gases but is heated by heating the walls of retort or box (containing the charge).

Mulliken Symbols. Terms applied in crystal field theory, e.g., A and B are singly degenerate electronic state, E doubly degenerate, T-triply degenerate, etc.

Mullite, Al₆Si₂O₁₃. Mineral aluminosilicate and a common constituent of intensely heated fireclay goods. Mullites increase the strength and refractness of bricks and resistant to corrosion.

Multiplet. A group of two or more closely grouped finer lines which make up the fine structure of the original line in a spectrum is known as a multiplet.

Multicentre Bond. A two electron bond formed by the overlapping of orbital from more than two atoms. In diborane (B₂H₆), the bridging is supposed to be by an sp³ hybrid overlap from each bridging is supposed to be by an sp³ hybrid overlap from each boron atom with the 1s orbital on the hydrogen atom. This multicentred bond is called a two-electron three-centre bond: The molecule is electron deficient.

Multidentate Ligand, Polydentate Ligand. A ligand with two or more sites at which it can coordinate. For example, H₂NCH₂ CH₂ NH₂.

Multiple Bond. Bonding between atoms in which there is overlap of orbitals in more than one position in space, e.g., a double

bond or triple bond. This additional bonding arises from overlap of atomic orbitals that are perpendicular to the inter-nuclear axis resulting in an increase in electron density above and below the inner nuclear axis. Such bonds are called π (pi) bonds. Generally, the primary bond is a σ (sigma) bond with extra

Multiple effect Evaporator. An evaporator containing several evaporator units in series. 1 kg of steam can evaporate. approximately, (N kg of water where N=the number of effects of steam generated from units of evaporator.

Multiple Proportion Law of (Dalton's Law). When two elements A and B combine to form more than one compound, the weights of B that combine with a fixed weight of A are in small whole number ratio. For example, in nitrous oxide (N₂O₄), nitric oxide (NO) and dinitrogen tetraoxide (N₂O₄), the amounts of nitrogen combined with a fixed weight of oxygen are in whole

Muon. A \u03c4-meson.

Muntz Metal. A brass-often referred as 60% "brass" which contains Cu (60%) and Zn (40%). Used in the manufacture of

Muonium. An electron+a muon.

Muriate. An obsolete name for a chloride, e.g., KCl.

Muriatic Acid. An obsolete name for hydrochloric acid.

Musacone, 3-Methylcyclopentadecanone, C₁₆H₃₀O.

Yellow liquid, B.P. 330°C. Obtained from the scent glands of

Musk, Artificial, C₁₁H₁₃M₃O₆. 2, 4, 6-Trinitro-3-ter-butyltoluene (musk bar). Yellow crystals M.P. 97°C. Used in perfumery. Musk Ketones. 4-tert-butyl 2, 6-dimethyl-3, 5-dinitroaceto-phe-

Musk Xylene. 1-ter-butyl-3, 5-dimethyl-2,4,6 trinitrobenzene.

Mustard Gas, 2,2'-dichlorodiethyl Sulphide, C₄H₈Cl₂S. Colourless oily liquid, M.P. 13-14°C, B.P. 215-217°C, (S₂Cl₂+ethene at 30-35°C). A powerful vesicant and poison which causes conjunctivities and temporary blindness. Decomposed by bleaching powder.

Mustard Oil. See allyl isothiocyanates.

Mutagens. Chemical substances or agents (physical) which increase the frequency of mutation above the normal level.

Mutarotation. A change in optical rotation of solutions with time. Optically active substances bring about this change which is catalysed by acids or bases.

Mutase. An enzyme.

Myogen. A protein of albumin class.

Myoglobin. The oxygen-carrying protein of mammalian muscle.

Myosin. An important protein of muscles belonging to globulin class of proteins.

Myrcene, C₁₀H₁₆. An alicyclic monoterpene found in some essential oils, e.g., in verbena oil and oil of hops.

$$\begin{array}{c} \text{H}_3\text{C} \\ \text{C=CH-CH}_2\text{-CH}_2\text{-C-CH=CH}_2 \\ \text{H}_3\text{C} \\ \end{array}$$

Myristic Acid, n-tetradecanoic Acid, C₁₄H₂₈O₂. CH₃ [CH₂]₁₂ COOH. M.P. 58°C, B.P. 250°C/100 mm. A fatty acid found in milk and vegetable oils.

Myrosin. An enzyme belonging to sulphatase class and found in black and white mustard seeds.

Myrtenal, C₁₀H₁₄O. An unsaturated aldehyde found in false camphor wood oil. B.P. 91.75/12.5 mm.

Myrtenol, C₁₀H₁₆O. A primary alcohol obtained from α-pinene.

N

N. Nitrogen.

Na. Sodium.

NAD. Nicotinamide adenine dinucleotide.

NADP. Nicotinamide adenine dinucleotide phosphate.

Nano, (n). A prefix denoting 10⁻⁷.

Napalm. A gel of gasoline obtained by adding soap of naphthaenic and palmitic acids. Used in military flame bombs and incendiary.

Naphtha. A coal tar product containing mostly of xylenes and higher homologues. Special boiling point spirits (SBP) with boiling range of about 40-150°C.

Naphthalene, C₁₀H₈. A white crystalline solid with a distinct smell of moth balls. Found in both the middle-and-heavy oil fractions of crude oil and is detained commercially from petroleum fractions by demethylation of methyl naphthalenes with H₂ at 750°C and 10-70 atom.

Gives additional and substitutional products more easily than benzene. Forms wax like compounds on exhaustive chlorination. Forms dyestuffs intermediates on nitration and sulphonation. Substitution in the ring forms two series namely, insecticides.

Naphthalene Ring System. Two or more than two benzene rings fused together to form a big structure.

Naphthalene Sodium, [C₁₀H₈]-Na⁺. A greensolution used as a metalation reagent. Prepared from Na metal plus naphthalene in some ethers like THF or diglyme.

Naphthalene Sulphonic Acid. Naphthalene monosulphonic acid (naphthalene+sulphonic acid) is always obtained as two isomers depending upon the temperature of the reaction which are:

Naphthalene-1-sulphouic acid (2H2O) a dihydrate, M.P. 90°C.

Naphthalene-2-sulphonic acid (3H₂O), a trihydrate, M.P. 83°C. Forms nitro derivatives.

Naphthalene disulphuric acids. Four isomeric forms (prolonged sulphonation of naphthalene), are obtained.

Naphthalene trisulphonic acids. Only 1,3,5 or 1,3,7-acids are prepared by drastic sulphonation of naphthalene or its mono or di-sulphonic acids). 1,3,6-acid is used for the preparation of H-acids.

Naphthalic Acid, 1,8-Naphthalene Dicarboxylic Acid.

Naphthalenes. Saturated hydrocarbons having at least one closed ring of carbon atoms (cyclo-alkanes), e.g., cyclohexane.

Naphthoic Acids. Carboxylic acids of napathalenes

1-Naphthol, or α -Naphthol, $C_{10}H_8O$,

Colourless crystalline, M.P. 94°C, B.P. 278°C, soluble in alcohol, ether, benzene, etc. Occurs in coal-tar fractions. Prepared by caustic soda fusion of sodium naphthalene-1-sulphonate or by high temperature alkaline digestion of 1-chloronaphthalene. Forms phthalic acid on oxidation. Forms a series of dyes on diazotization. Forms picrate (M.P. 190°C).

2-Naphthol or β-naphthol, C₁₀H₈O,

White crystals, M.P. 122°C, B.P. 285-286°C, soluble in C₆H₆, CHCl₃, C₂H₅OH, etc. Occurs in traces in high boiling coal-tar fraction. Prepared by caustic soda fusion of sodium naphthalene-2-sulphonate. Oxidized to phthalic acid, chlorinated and sulphonated. An antiseptic that forms a characteristic yellow colour with bleaching powder in aqueous solutions coupled with diazotized base to form a series of dyes.

1-Naphthylamine or α-Naphthylamine, C₁₀H₉N,

Colourless crystals, M.P. 50°C, B.P. 301°C, soluble in alcohol and ether. Basic and forms soluble salts with mineral acids. Prepared from ammonia plus α-naphthol (under pressure) or by reducing 1-nitronaphthalene with Iron/HCl. Diazotised easily and used in so many monoazodyes. Forms naphthionic acid on sulphonation.

2-Naphthylamine or β-Naphthylamine, C,0H₉N,

Lustrours leaflets, M.P. 112°C, B.P. 294°C, soluble in hot water and alcohol. Forms hydrochloride and sulphates. Prepared

by Bucherer's method (heat β -naphthol+strong solution of ammonium sulphite and ammonia at 150° for 8 hours). Used as an end component in azo dyes.

O.CO.NHCH₃

1-Naphthyl-N-methyl Carbamate, $C_{12}H_{11}NO_2$,

, (Sevin).

White soiid, M.P. 142°C. Prepared from 1-naphthol, plus methyl isocyanate or PH₃ with a base. A contact insecticide.

Naphthol Yellow S. A nitro-dye. Used for dyeing the animal textile fibres, chiefly wool and for colouring food (non-poisonous).

Narcotine, C₂₂H₂₃O₇N. Colourless needles, soluble in CHCl₃. An alkaloid of opium. Acts medically as that quinine.

Nascent Hydrogen. An active form of hydrogen produced by electrolysis or chemical reaction and used in situ for reduction. Used to produce the hydrides of P, As and Sb, which are not easily formed by oridinary hydrogen. The powerful reducing action may be due to reaction at surface or hydride species.

Natta Process. A process for the manufacture of isotactic poly-

Natrolite, Na₂ (Al₂ Si₃ O₁₀), 2H₂O.

Natural Gas. Gases which originate from underground accumulation which may or may not be incorporated with crude oil. It contains mainly 95% methane and other low boiling hydropresent (associated with N₂ and CO₂ only (dry gas). If oil is of higher hydrocarbons such as pentane, hexane, etc., and is known as wet gas. Used as a fuel.

Nb. Niobium.

Ne. Neon.

Neel Point. The temperature at which magnetic susceptibility

Negative Adsorption. When add reption takes place at a surface from a solution one component may be adsorbed more on the surface rather than in the bulk and simultaneously the other the bulk. The latter adsorbed less on the surface rather than in on the surface. It is governed by Gibb's adsorption equation which states that any solute which increases the interfacial tension of the system will be negatively adsorbed.

Neodymium, Nd. At. No. 60, At. wt. 144'24, M.P. 1010°C, B.P. 3127, D 7'00. A typical toxic silvery element of lanthanide series. Occurs in association with other lanthanides. Forms compounds in +3 state. Used in glasses and in capacitors.

Neon, Ne. At. No. 10, At. wt. 20'179, M.P. -248'67, B.P. -246'05. A noble gas of rare gas series with electronic configuration 1s², 2s² p⁶. Occurs in air (1'8×10⁻³ by volume of air) and can be obtained by fractionation of liquid air. Used in fluorescent tubes (red neon lights), electrical equipments, in Gas lasers, etc.

Neopentyl. The group Me₃C CH₂-.

Neophyl. The group Ph Me₂C CH₂-.

Neoprene, Polychloroprene. A type of synthetic rubber formed by polymerization of 2-chlorobuta-1, 2 diene (H₂C=CH-CCl=CH₂). The molecular weight is controlled by the addition of S or sulphur containing compounds during emulsion polymerization. It is resistant to solvents, to abrasion and has high tensality. Used in automove goods, as a fibre binder and in road and building construction.

road and building constraints
$$CH_2 = CH - C = CH_2 - \cdots \longrightarrow \begin{bmatrix} CH_2 - CH = C - CH_2 \\ CI \end{bmatrix} CI$$

$$CH_2 - CH = C - CH_2$$

$$CH_2 - CH = C - CH_2$$

$$CI$$
Neoprene rubber

Also used in making hoses for petrol and oil and containers for corrosive chemicals.

Nephelauxetic Effect. The change in electronic spectra of complexes due to varying degree of covalent character in the metal ligand bonds. Measuring parameter is β .

Nephelometry. A quantitative method used for the analysis involving the spectrophotometric estimation of the scattering of light by a colloidal suspension of a precipitate.

Neptunium, Np. At. No. 93, At. wt. 273'0482, M.P. 6403°C, B.P. 3902°C, D. 20'45. A toxic radioactive element of the actinide series of metals that was the first transuranic element to be synthesised. Occurs on earth in traces only in uranium compounds, formed by neutron capture of natural uranium. Prepared artificially, by neutron irradiation of ²³⁸U or ²³⁵U. Synthesised by reducing NPF₃ by Li (six crystalline forms). Separated by selective oxidation and solvent extraction method. Used to form ²³⁸Pu which is a major power source.

Np shows a strongly oxidizing +7-state and forms compounds similar to uranium. Electronic configuration is 5f⁵, 6s² p⁶, 7s²,

The metal dissolves in acid to form Np3+ oxidized by air to Np4+ and by mild oxidants to NpO2+. Forms halides such as NpF4 (orange), NpF5, NpCl4, NpBr4.

Nernst Equation. An equation corelating the electromotive force of a cell and concentrations or, more accurately, the activities of the reactants and products of the cell reaction. For a reaction of a galvnic cell

the electromotive force is given by

$$E=E^{\circ}-RT\ln \left(\frac{ac^{\circ}.\ aD^{d}}{a^{a}A.\ a^{b}B}\right)$$

where a, b and c, d are the number of moles of reactants and products respectively, a_n denotes the activities of reactants and products under given the conditions and E° is the standard electromotive force of the cell.

Nerol, C₁₀H₁₈O, (3,7-dimethyl·2, 6-Octadiene·l·ol). B.P. 225-226°C. A terpenic alcohol with the constitutional formula

A constituent of neroli, petit grain and of many essential oils.

Used as a constituent of perfume.

Nerolidol, C₁₅H₂₆O. B.P. 276°C. An alcohol found in neroli oil (+form).

Nerve Gas. Fluorophosphoric acid derivatives, e.g., esters obtained from Ag₂PO₃ and RX, are quite toxic and inhibit cholinesterase posed as nerve gas.

Nervone, C₄₈H₉₁NO₈. M.P. 180°C. A cerebroside.

Nervonic Acid, C₂₀H₄₆O₂, CH₃ [CH₂]₇CH=CH. [CH₂]₁₃ COOH. M.P. 43°C. A powder obtained from nervone.

Nessler's Reagent. An alkaline solution of mercuric iodide (HgI2) in potassium iodide (KI) used for detecting and estimating ammonia. Forms a brown colour or precipitate of NH2Hg2OI.

- Nessler's Tubes. Cylinders used for comparing turbidities and colours of solutions.
- Neuraminie Acid, C9H17NO8. An amino sugar present in animal tissues.
- Neurine, Trimethylvinyl Ammonium Hydroxide, C5H13NO. A liquid forming a crystallive trihydrate. Present in free and combined state in brain.
- Neutralization. The process of neutralization of an acid by a base or vice-versa. (H++OH-=H2O)
- Neutralization, Heat of. The amount of heat evolved when 1 gram equivalent of an acid is neutralized by 1 gram equivalent of the base. For example, the heat of neutralization for a strong acid and strong base in dilute solution is 13.7 Kcal (57.35 kJ) and the only reaction which takes place is $H^++OH^- \longrightarrow H_0O$.

The heat of neutralization depends upon the nature of the acid and base.

- Neutrino. An elementary particle without charge but with spin 1 and assumed to account for the conservation of angular momentum-in nuclear transformation. It seems to have variable finite mass and to oscillate from type to another.
- Neutron. A fundamental particle of atom, without charge (electrical) but mass equal to that of a proton. It has magnetic moment 1.91 B.M. and interact with atoms with unpaired electrons. Decays to form an electron and proton.
- Neutron Number, N. The number of neutrons in the nuclei of an atom and is equal to nucleon number (A) minus the proton number (2).
- Neville-Winther Acid. 4-Hydroxy-1-naphthalene sulphonic acid.
- Newland's Law (of Octave). When the elements are arranged in order of increasing atomic weight, the eighth element starting order of increasing atomic from a given one is the repetition of the first like the eighth note of an octave in music. For example, Li, has similarity to Na,
- Newton, N. The SI unit of force equal to the force required to accelerate 1 kilogram by 1 meter second⁻². (1N=1 kg ms⁻²)
- Ni. Nickel.
- An alloy of nickel containing Ni (60%), Fe(25%), Nichrome. Cr (15%).
- At. No. 28, At. wt. 58.69, M.P. 1453°C, B.P. 2732°C, D 8.90. A transition element of group VIII having electronic Nickel, Ni.

configuration 3d8, 4s2. Occurs in nature as mixed metal sulphide ores (e.g., millerite), as mixed arsenides, antimonides, sulphides, as garnierite and as pyrrhotite (Fen S_{n+1}). The ore is roasted to NiO and is reduced to Ni with C.

$$\begin{array}{ccc} 2 \text{ NiS+3O}_2 & \rightarrow 2 \text{NiO+2SO}_2 \\ \text{NiO+H}_2 & \rightarrow \text{Ni+H}_2 \text{O} \end{array}$$

The nickel is separated in volatilizers by converting it to nickel

carbonyl [Ni+CO ———→ Ni (CO)₄] followed by conversion to Ni (pure) at 180°C (Mond's process).

 $Ni(CO)_4 \rightarrow Ni+4CO$. volatile nickel carbonyl

Ni is silvery-white metal and is not tarnished by air but attacked by acids (except conc. HNO₃). It is not attacked by fluorine. Used as a catalyst in hydrogenation and in glass

Nickel Accumulator. A source of current with a discharge voltage about 1'2 volts. The reaction of the accumulator is

 $Fe(S)+NiO.OH+H_2O \rightarrow Fe(OH)_2(S)+Ni(OH)_2$

On charging, the Fe(OH)2 is reduced to Fe and the Ni (OH)2 oxidized to Ni₂O₃. It has a low current efficiency and more expensive as compared to lead accumulators. It has low weight

- Nickel Alloys. They are anticorrosive and expensive, e.g., Monel metal (Cu 30%, Ni 70%), Constantan (Cu 60%, Ni 40%), German silver (Cu 55%, Ni 20%, Zn 25%), Platinoid (Cu 60%, Ni 14%, W 2%), invar, nilo, etc.
- Nickel Ammonium Sulphate, Ni(NH₄)₂ (SO₄)₂, 6H₂O. Blue-green obtained from nickel sulphate and ammonium sulphate. Used
- Nickel Arsenide, NiAs. This compound, NiAs, has given its name to an important crystal structure type. The structure is hexagonal to NiS. Many elements crystallize with this structure,
- Nickelates. Compounds such as nickelates (IV) and nickelates (III),
- Nickel Bromide, NiBr₂. Prepared in anhydrous and hydrous forms, e.g., NiBr₂, 3H₂O; NiBr₂, 6H₂O and NiBr₂, 9H₂O.
- Nickel Carbonate. Unstable carbonates, e.g., NiCO₈, 6H₂O and

- Nickel Carbonyl, Ni(CO)4. A colourless liquid with high toxic vapours. Obtained by passing CO over finally divided Ni at 60°C (Ni+4CO→Ni(CO)₄ followed by fractional distillation. Decomposes to pure nickel at 180°C.
- Nickel Chloride, NiCl2. A yellow coloured anhydrous chloride obtained by heating nickel powder or dust in chlorine (fire is produced). Forms hydrates with octahedral nickel. A hexahydrate NiCl₂, 6H₂O is formed from its solution which forms a double salt (NiCl₂.NH₄Cl.6H₂O) with ammonia chloride and a violet compound NiCl2, 6NH3 with ammonia.
- Nickel Dimethyl Glyoxime, C₈H₁₄N₄NiO₄. A substance formed during the estimation of nickel and which is finally weighed.
- Nickel Fluorides, NiF2. Yellow green forms perovskites, e.g., KNiF₃, K₃NiF₆ and K₉NiF₆.
- Nickel Hydroxide, Ni(OH)2. Apple green and used as an oxidizing agent.
- Nickel Iodide, NiI2. Forms hexahydrate.
- Nickel Nitrate, Ni(NO₃)₂, 6H₂O. Green crystals (Ni+N₂O₄).
- Nickel Organic Derivatives. Forms bis-n-allyls, e.g., (h3-C3H5)2Ni —a catalyst for cyclotelomerization of diolefins.
- Nickel Oxide, NiO (Nickelous Oxide). A pale green powder (heat Ni(OH)₃. NiCO₃ or NiNO₃ in absence of air). Dissolves in dilute acids to form green solutions of nickel salts.
- Nickel Oxide, Ni₂O₃ (Nickelic Oxide). A black powder (heat NiO in air). Exists as dihydrate, Ni₂O₃, 2H₂O.
- Nickel Sulphate, NiSO4, xH2O. A soluble salt which forms double sulphates Ni(NH₄)₂SO₄, 6H₂O.
- Nickel Vitriol. NiSO4, 7H2O.
- Nicol Prism. A prism cut from a calcite crystal and is used to polarize the light i.e., plane spolarized light obtained from plane
- Nicotinamide Agenine Dinucleotide (NAD). An important coenzyme which functions as a hydrogen transfer agent. It was formerly known as DPN. It is reduced to NADH. Obtained biosynthetically from nicotinamide or nicotinic acid.
- Nicotinamide Adenine Dinucleotide Phosphate (NADP). An enzyme of pyridine nucleotide group. It was also known as TPN (triphospnopyridine nucleotide).
- Nicotine, 3-(1-methyl-2-pyrrolidyl) Pyridine. B.P. 246°C. A strong, base, laevorotatory liquid with pyridine like smell. Muscible

$$H_2C$$
 H_2C
 CH
 $=$
 N

CH, with water below 60°C and above 210°C. Soluble in alcohol and ether. Occurs in tobacco plants combined with citric and malic acids. An alkaloid, used as an insecticide.

Nicotinic Acid, Niacin, 3-Pyridine Carboxylic Acid, C6H5NO2. M.P. 232°C. An essential component of mammlian diet. Manufactured from pyridine or quinoline. Forms NAD (nicotinamide adenine dinucleotide).

Nido. A term applied to open structures.

Ninhydrin (C₉H₄O₃, 3H₂O). A colourless organic compound that forms a blue colour with ammonia. Used for detection of proteins peptides and amino-acids (forms blue colour on heating with ninhydrin). Light brown crystals which lose water at 120-130°C.

Used as a spray reagent in paper chromatography.

Niobium, Nb. At. No. 41, At. wt. 92.9064, M.P. 2468°C, B.P. 4742°C, D 8.57. A transition element of group V with electronic configuration $5s^1 4d^4$ also known as columbium, Cb. Chief mineral is columbite tantalite (Fe, Mn) (Nb, Ta)₂O₆. Obtained by reduction of Nb₂O₅ with carbon. Dissolves in HNO₃-HF. Important oxidation state is +5. Forms hydrides, borides (NbB, Nb₃B₂), carbides (Nb₄C₃, NbC) and nitrides (Nb₂N, Nb₄N₅, Nb₅N₆). Also forms sulphides (Nb₃S₄ and NbS₂) and some complexes. Used in welding, special steels and nuclear reactor work.

Niobium Organometallic Compounds. Nb forms cyclopentadienyl

Niobium Oxides. Niobium forms oxides such as Nb₂O₅ (dense, white inert material) which forms niobates (KNbO₈). Prepared by hydrolysis of niobium halides followed by dehydration.

Witramide, H.N.NO.

- Nitrates. A sal of nitric acid (HNO₃). All metal nitrates are soluble in water. Nitrates are nested by the 'ring test' (Nitrate solution +ferrous sulphate solution+conc. H₂SO₄ from the sides of the tube—formation of a brown ring at the junction of liquids). Nitrates of metals such as Na, K, Ca one of commercial importance. Organic nitro compounds or ammonium nitrate constitute a part of explosives. Also used as a fertilizer.
- Nitrating Agent (acid). A (1:1) mixture of conc. HNO3 or conc. H₂SO₄ used for nitration, e.g., the formation of 2, 4, 6-trinitrotoluene (TNT) from toluene which is an explosive. The attacking species is NO2+(electrophile substitution).
- Nitrene Group, (RN=.) (the amine group). Nitrene derivatives are formed by condensation, e.g., [WF5NMe].
- Nitric Acid, HNO₃ M.P. -42°C, B.P. 83°C (decomposes). A colourless fuming corrosive liquid prepared in the laboratory by an alkali metal nitrate and sulphuric acid (NaNO₃+H₂ SO₄→ NaHSO₄+HNO₃). Commercially prepared by the catalytic oxidation of ammonia and is supplied as a concentrated nitric oxidation of ammonia and is supplied as a concentrated nitric acid (68% of the acid, D 1.4) which is generally yellow due to dissolved oxides of nitrogen. Pure HNO₃ is obtained by redistilling the commercial acid under reduced pressure (98%,

A strong oxidizing agent and oxidizes non-metals to oxyacids, metals to the corresponding nitrates plus oxides of nitrogen.

Reactions of non-metals

$$S+6HNO_3=H_2SO_4+6NO_2+2H_2O$$

 $I_2+10HNO_3=2HIO_3+10NO_2+4H_2O$
 $P+5HNO_3=H_3PO_4+5NO_2+H_2O$

Oxidizes amorphous C to mellitic acid, C3 (COOH)6 and finally to CO₂. Graphite forms a yellow insoluble acid, graphitic acid $(C_{11}H_4O_6).$

Reactions of metals

Mg+2HNO₃
$$\rightarrow$$
Mg (NO₃)₂+H₂
Cu+4HNO₃ \rightarrow Cu (NO₃)₂+2NO₂+2H₂O
3Ag+4HNO₃ \rightarrow 3Ag NO₃+NO+2H₂O
6Hg+8HNO₃ \rightarrow 3Hg₂ (NO₃)₂+2NO+9H₂O
dilute
3Hg+8HNO₃ \rightarrow 3Hg (NO₃)₂+2NO+4H₂O
Concentrated

Conc. nitric acid (1 part)+conc. HCl (3 parts) form aqua regia, a very good solvent for noble metals such as Au, I't, etc. Used in the production of fertilizers (NH₄ NO₃), explosives, dyestuffs, etc.

- Nitric Oxide, Nitrogen Monoxide (NO). A colourless gas, insoluble in water but soluble in solutions containing Fe2+ ions due to the formation of (Fe NO)2+ (the NO can be recovered by heating the solution). Prepared from nitric acid plus copper turnings. Commercially prepared by the catalytic oxidation of ammonia or direction combination of N_2 and O_2 in an electric arc $(N_2 + 1)$ $O_2\rightarrow 2NO$). It decomposes above 1000°C to form the parent elements. Combines immediately with O_2 to form nitrogendioxide $[NO(g)+O_2(g)\rightarrow 2NO_2(g)]$.
- Nitrides. Compound of nitrogen and other elements. Prepared from element plus N2 or NH8. Two types of nitrides, e.g., Li₃N (obtained from electropositive elements) and nitrides of transition metals. Covalent nitrides are formed by less electro-positive elements. Used in ceramics.
- Nitriding. A process of surface hardening steel (in NH₃) with the
- Nitride Complexes. Complexes with N3 ligand.
- Nitrile, (cyanide). A type of organic compound containing the -CD groups. Colourless liquids undergo hydrolysis and reduction. Used as fuel additives, weed control, lubricants, etc.
- Nitrile Rubber. A copolymer of butadiene and acrylonitrile an oil and solvent resistant rubber which is used in gasketes, seal-
- Nitrites. Salts of nitrous acid containing [NO₂] ion. Alkali nitrites are used in bacons as preservative.
- Nitroamines. Organic compounds having both -NO₂ (nitro) and
- m-Nitroaniline, 1-Amino-3-Nitrobenzene, $C_6H_6N_2O_2$,

Yellow needles, M.P. 114°C, B.P. 285°C. Used

in the preparation of 930 dyes.

p-Nitroaniline, 1-Amino-4-Nitroaniline, C6H6N2O2,

Yellow crystals, M.P. 147°C.

Nitroanthraquinones. Nitrated anthraquinones (at o-pasition).

NO. Colourless, refractive liquid Nitrobenzene, C₆H₅NO₂,

with characteristic smell, B.P. 209°C, D 1'209. Miscible with alcohol ether and benzene, easily volatile in steam. Prepared by the action of conc, HNO3 and conc. H2SO4 on benzene at 40°-60°C.

$$\bigcirc + \text{ Conc. HNO}_{8} \xrightarrow{\text{Conc. H}_{2}\text{SO}_{4}} \rightarrow \bigcirc + \text{ H}_{2}\text{O. Further}$$

nitration gives m-dinitrobenzene, sulphonation gives m-nitrobenzene sulphonic acid. On reduction it gives azoxybenzene, then azo benzene and aniline depending upon the condition.

$$\frac{\text{Zn/NaOH}}{2\text{C}_6\text{H}_5\text{-NO}_2 + 6[\text{H}]} \xrightarrow{\text{C}_6\text{H}_5} \text{C}_6\text{H}_5 - \text{NH} - \text{NH} - \text{C}_6\text{H}_5} \text{C}_6\text{H}_5 - \text{NH} - \text{NH} - \text{C}_6\text{H}_5} \text{NH} + \text{NH} - \text{NH} - \text{C}_6\text{H}_5} \text{NH} + \text{NH} - \text{NH} - \text{NH} - \text{NH} - \text{C}_6\text{H}_5} \text{NH} + \text{NH} - \text{NH}$$

Used as an important dyestuff intermediate being used as a first component of azo dyes.

Nitro Compounds. A type of organic compounds containing the nitro group (-NO₂) attached to an aromatic ring. Prepared by nitration. (Conc. H₂ SO₄+Conc. HNO₃)

Nitro Compound

Nitro-dyes. The coloured salts of several nitrophenols or derivatives of sulphonic acid, e.g., picric acid

Nitrogen, N. At. No. 7, At. wt. 14:006, M.P. -209:86°C, B.P. rugen, N. At. No. 1, Ment of the V group, colourless gas ex-- 195.8°C. The first element of the Present in air (75.5% by isting as N₂ and very electronegative. Present in air (75.5% by weight), in proteins and also as nitrates and ammonia. Electronic configuration) $2s^2p^3$. A typical non-metal forming typical polarized covalent bond. Commercially prepared by fractional distillation of liquid air and also prepared (in the laboratory) by heating NH₄ NO₂ (Na NO₂+NH₄ Cl). With electropositive elements N³⁻ ion may be formed. Also combines with O₂, H₂ (Haber's process), CaC₂ (to form calcium cyanamide), etc. Active nitrogen is produced by electric discharge (excited H₂). Pure nitrogen is prepared by thermal decomposition of azides:

$2NaN_3 \rightarrow 2Na + 3N_2$

Forms oxides such as N_2O (nitric oxide), N_2O_3 (nitrogen sesqoxide, an unstable oxide), NO_2 (nitrogen dioxide), N_2O_6 (nitrogen pentaoxide) and N_2O_4 (nitrogen tetraoxide).

Nitrogen has two isotopes ¹⁴N, the common isotope and ¹⁵N (about 0.3%). ¹⁵N isotope is used as a marker in mass spectroscopy. Nitrogen is used in fertilizers and liquid nitrogen is used as a refrigerant.

- Nitrogenase. A complex of metal containing proteins present in nitrogen-fixing bacteria and blue-green algae.
- Nitrogen Chlorides. Nitrogen forms nitrogen trichloride, NCl₃, an explosive oily liquid (Cl₂+NH₄Cl solution). Also binary halides such as NF₃, NCl₃ and halogen azides, ClN₃ are also known. Other nitrogen halides are NHCl₂, NH₂Cl, NF₂Cl and NFCl₂.
- Nitrogen Dioxide, NO₂. A brown gas obtained from dinitrogen tetraoxide at 140° C (N₂O₄ \longrightarrow 2NO₂). Further heating forms NO and O₂ (2NO₂(g) \longrightarrow 2NO(g)+O₂(g).
- Nitrogen Fixation. The use of atmospheric nitrogen to form compounds of commercial importance such as nitric oxides (conaformate, bicarbonate and sulphate), ammonia (used as (calcium cyanamide and nitrides). Bacteria present in the roots free living soil-organism also fixes nitrogen in nature.
- Nitrogen Fluorides, NF₃ (nitrogen trifluoride). A very in ert substance (fused ammonium hydrogen chloride is electrolysed). Cisand trans-difluorodizine (FN=NF) and F₂N.NF₂ are also known.
- Nitrogen Hydrides. Nitrogen and hydrogen compounds such as, NH₃ and other derivatives N₂H₄, etc.
- Nitrogen tri-iodide, NI₃. A black powder (NH₃+KIBr₂).
- Nitrogen Mustards. Some poly-(β-chloroethyl) amines with general formula RN(CH₂CH₂Cl)₂ where R is an alkyl, alkylamine of

alkyl chloride group. Resembles mustard gas in properties. Used in the treatment of Hodgkin's disease.

Nitrogen Oxide. See nitric oxide.

Nitrogen Oxyacids

Hyponitrous Acid, $H_2N_2O_2$. Its alkali salt solution is obtained from KNO₂ and Na amalgam; $Ag_2N_2O_2$ is precipitated from solution and $H_2N_2O_2$ is formed from Ag_2 N_2O_2 and cold dilute HCl in ether. The $H_2N_2O_2$ is filtered off.

$$Ag_2N_2O_2+2HCl\rightarrow H_2N_2O_2+2AgCl$$

The hyponitrite ion is trans (O-N-N-O)2-.

Nitric Acid, HNO, See nitric acid.

Nitrous Acid, HNO₂. Its free state existence is doubtful and it is likely that when N_2O_3 dissolves in water at the ice temperature, a blue liquid thus formed consists mainly of nitrous acid. Highly unstable and decomposes:

$$N_2O_3+H_2O\rightarrow 2HNO_2$$
 (at 0°C)
 $2HNO_2\rightleftharpoons HNO_3+2NO+H_2O$ (at low temperatures)
 $2HNO_2\rightleftharpoons NO+NO_2+H_2O$ (at high temperatures)

Acts as a reducing agent, e.g., H_2O_2 is reduced to water and Br_2 water is reduced to hydrobromic acid (HBr). Acts as an oxidising agent, e.g., liberates iodine from iodides $(2HI+2HNO_2 \rightarrow 2NO+2H_2O+I_2)$, H_2S is oxidized to S and SO_2 is oxidized to H_2SO_4 .

The probable structure is:

$$H-N_{H}^{O} \rightleftharpoons HO-N=0$$

Hyponitric acid, H₂N₂O₃. Trioxy dinitrate. The free acid is unstable.

Nitroglycerine, Glyceryl Trinitrate, C₃H₅N₃O₈ CH₂(ONO₂). CH₂(ONO₂). A highly explosive oily liquid prepared CH(ONO₂). CH₂(ONO₂). A highly explosive oily liquid prepared by reacting with a mixture of nitric and sulphuric acids. In pure form, it is colourless, odourless, insoluble in water with pure form, it is a nitrate ester and is highly explosive. It is M.P. 8°C. It is a nitrate ester and is highly explosive. It is never used alone due to high sensitivity. Used as a pain relief never used alone due to high sensitivity. Used as a pain relief or chest pain in angina. Also used in dynamites and propellants.

Nitron, C₂₀H₁₆N₄. Used as a precipitating agent for NO₃⁻, ClO₄⁻, PF₆⁻ and WO₄²⁻ (in ethanoic acid solutions).

1-Nitronaphthalene, C₁₀H₇NO₂, (NO₂). Yellow crystal, M.P.

61°C, B.P. 304°C. Prepared by nitrating naphthalene. Used in the preparation of 1-naphthyl-amine.

Nitronium Ion. NO2+ ion.

Nitronium Salts. Salts having NO2 ion. Prepared by reacting strong acids and conc. HNO₃ or NO₂ plus oxidizing agent.

Nitroparaffins, C_nH_{2ⁿ+1}NO₂. Colourless toxic liquids, sparingly soluble in water. Prepared from AgNO₃ plus alkyl halide or by reacting propane and HNO₃ in vapour phase at 400°C (lower nitro-paraffins). Forms amine when reduced with Sn and HCl. Lower paraffins are used as solvents, propellants, etc.

Nitrophenols, CoH4 (OH) (NO2). Organic [compounds formed by direct or indirect nitration of phenols. Three isomers, of nitrophenols are as follows:

(i) 2-nitrophenol or o-nitrophenol, (). M.P. 45°C. Bright yellow solid (aitration of phenol). Used in dyestuffs and photo-

(ii) 4-nitrophenol or p-nitrophenol, (O), M.P. 114°C. A

white crystalline solid.

Nitroprussides. Complexes containing the [Fe(CN)₅NO)]²⁻ ion and having a formula M₂ [Fe(CN)₅NO]. For example, sodium nitroprusside, Na₂[Fe(CN)₅(NO)], 2H₂O which is used as a test for soluble sulphides, with which it gives violet colour.

Nitrosoamine Yellow oils or solids containing (-NO) group and with a general formula NO. Obtained from HNO₂+

Nitroso Dyes. Dyestuffs having nitroso group (-NO), e.g., quinone

Nitrosomethyl Urea, NH₂CON(NO)CH₃. A colourless solid, M.P. 73°C. Prepared from NaNO₂ plus methyl urea nitrate (in solution).

Nitrosonium Hydrogen Sulphate, Chamber Crystals, NaHSO₄. M.P. 73°C. A white solid used in diazotization.

Nitrosonium Salts. Salts containing NO+ ion.

4-Nitrosophenol, C₆H₅NO₂. Greyish brown crystals, M.P. 124°C (decomposes) Prepared from HNO₂ plus phenol. Used to form indophenols (with p-nitrosophenol) which are used for the commercial preparation of sulphur colours.

Nitrososulphuric Acid, Nitrosulphuric Acid, Nitro Sulphonic Acid. Alternate names for nitrosyl sulphuric acid, NOHSO₄.

Nitrous Acid. See nitrogen oxy acids.

Nitrous Fumes. A mixture of NO and N2O4 used for oxidation.

Nitrous Oxide. See nitrogen oxides.

Nitryl Ion (nitronium ion). The positive ion NO2+.

Nitryl Halides, XNO₂ (where X=F or Cl). Halogenated nitro derivatives, e.g., FNO₂ and ClNO₂.

n.m.r. Nuclear magnetic resonance.

n.m.r. Shift Reagents. Substances used to simplify n.m.r. spectra, e.g., europium complexes.

No, Nobelium.

Nobelium, No. At. No. 102. The metal has not been prepared. It shows +2 and +3 oxidation states. No solid compound is known. 235No is formed by the action of 12°C and 13°C nuclei on a cerium target.

Noble Gases, (rare gases, 0 group elements). The elements helium, neon, argon, krypton, xenon and radon. Occupy position between the highly electronegative group VII elements and the highly electropositive group I elements. They have all completely filled s and p with the electronic configuration s^2p^6 (excep He, s^2). All occur as minor constituents of the atmosphere. He is a component of some natural hydrocarbons whereas radon is obtained from radium emanations. Separated by fractional distillation. Used to provide inert atmosphere, e.g., welding (Ar), filling light tubes (Ne) and bulbs (Ar), gas stirring in high temperature metallurgy (Ar). Liquid He is used in cryoscopy Xe, Kr and Rn form some compounds.

Node. A region of very low electron density in an atom or molecule (wave equation).

- Non-Aqueous Solution. A solution prepared from solvents (e.g., C₂H₅OH, CHCl₃, NH₃, BrF₃, etc.), other than water.
- Non-aqueous Solvents. Solvents other than water.
- Non polar Molecules. Molecules which have zero dipole moment, e.g., symmetrical non-polar compounds (covalent compounds) such as CCl₄.
- Non-stoichiometric Compounds. Compounds in which the ratios of number of atoms present do not follow simple integers, e.g., silicates.
- Nonylphenol, C₁₅H₂₀O. A phenol used to produce phenolic resinsand surfactants.
- **nor.** A prefix indicating the loss of a $-CH_2$ group in an organic compound. For example, nor-nicotine $(C_{10}H_{14}N_2)$ —nor-nicotine $(C_9H_{12}N_2)$. When applied to terpenes, it means loss of all $-CH_2$ groups from parent compound. Also applied in amino acids.
- Normality, N. The number of gram equivalent of solute per litre of solution. IN solution of oxalic acid (hydrous acid) contains 63 grams of the acid in 1 litre of solution (eq. wt. of oxalic acid is 126/2, i.e., 63).
- Normal Temperature and Pressure, (NTP) or (STP).

Np. Neptunium.

Nuclear Magnetic Resonance (NMR). A method for studying nuclear spins (or magnetic moments). The substance to be investigated is placed in a very strong, uniform and steady field. This resultsin 'splitting' of the nuclear energy levels. The sample is then placed in an additional, weak and oscillating magnetic field whose frequency is 'scanned' over an appropriate range. At precise frequencies the nuclear nate with the field in undergoing transition between the magnetic energy levels. The resonance is amplified and recorded (using search coils). Hydrogen nuclei, for instance, can have two energy states, and the transition between the two occur by absorption of the radio frequency radiation. This technique is used to study molecular structures because the resonance frequencies of a specific element are influenced by the electronic snvironment in which the atom is situated, i.e., the position of the atom in the molecule. For example, in ethanol there are three frequencies corresponding to hydrogen atoms on the CH₃, the CH₂ and the OH groups. The intensity of absorption also depends on the number of hydrogen atoms (31 2:1). Sometimes nuclei couple together to give fine structures in the NMR spectrum and the extent and nature of the coupling. provides further information on molecular structures.

Nuclear Paramagnetism. The total angular momentum of the spin of several nuclei is given by

$$\frac{h}{2\pi}\sqrt{1(I+1)}$$

where I=the nuclear spin number $\left(I=0, \frac{1}{2}-, 1, \frac{3}{2}\cdots\right)$

and h=the Planck's constant, I=0, for non-spinning nuclei. This technique is used to study nuclei having I greater than 0.

Nuclear Quadrupole Moment. The electrical quadrupole moment of the nucleus which is used to study molecular structures (NQR),

Nuclear Spin. Nucleus has a property of spin and the angular moment of particles is expressed in units of $\left(\frac{h}{2\pi}\right)$.

Nuclease. An enzyme which hydrolyses nucleic acids.

Nucleic Acids. The part of nucleo-proteins and essential constituents of all living cells as they provide the necessary hereditary information enabling high specific proteins to be constructed. There are two types of nucleic acids.

Deoxyribose Nucleic Acid (DNA). Found in nuclei of cells. It is built of the nucleotides of the purines adenine and guanine and the pyrimidines cytosine and thymine. A molecule, consists of two helical chains coiled round the same axis, each chain comprising of alternate phosphate and sugar groups linked together through carbon 3 and 5 of the pentose molecule, and with the bases attached to the pentose. The chains are held together by hydrogen bounds between the bases. The two linked bases must be a purine and other a pyrimidine.

Ribose Nucleic Acid. Found in the cytoplasm. In RNA the sugar is ribose and uracil replaces thymine. Various types of RNA depending upon their molecular weight are known. For example, messenger RNA conveys information in the form example, messenger RNA conveys information in the form of the sequence of bases from DNA to RNA on the ribosomes, and transfer or soluble RNAs which are specific carrier molecules for amino-acids during protein synthesis on ribosomes with ribosomal RNA as the template.

Nucleon. A name for particles having mass number 1, i.e., proton + neutron.

Nucleon Number (mass number), (A). The number of nucleons in an atom, i.e., the number of protons + number of neutrons in an atomic nucleus.

Nucleophile (nucleophilic reagent). An electron rich ion or molecule that takes part in an organic reaction, e.g., the negative ions (Br-, CN-) or a molecule with a lone pair of electrons (NH₃, H₂O). Nucleophile always attacks on the positively charged part of the molecule Groups which act by donating or sharing their electrons, e.g., OH ion, OR and SR groups.

Nucleophilic Addition. Addition of a small molecule to an unsaturated organic compound, e.g., the carbonyl compounds undergo nucleophilic additions. Carbonyl group (>C=O) is constituted by a σ bond and a π bond. Oxygen being more electronegative than carbon, there is high electron density on oxygen than that on carbon. The carbonyl group is somewhat having the following structures

$$>C=O$$
 \leftrightarrow $>C=O$ or $>C=O$

Nucleophiles always attack at the positive centre of the molecule. The addition of hydrogen cyanide to propanone forms acetone

CH₃ C=0
$$\xrightarrow{-E, -M}$$
 CH₃ C $\xrightarrow{-\overline{O}}$ $\xrightarrow{\overline{CN}}$ CH₃ CH₃ C $\xrightarrow{\overline{CN}}$ CH₃ CH₄ CN CH₃ CH₃ CH₃ CH₄ CN CH₃ CH₄ CH₄ CN CH₃ CH₄ CN CH₄ CH₄ CN CH₃ CH₄ CH₄ CN CH₄ CN CH₄ CH₄ CN CH₄ CH₄ CN CH₄ CH₄

Nucleophilic Substitution (S_N) . A reaction involving the substitution of an atom or group of atoms in an organic compound with a nucleophile as the attacking substituent. Nucleophilic substitution occurs in compounds in which a strongly electronegative atom or group leads to a dipolar bond. For example, hydrolysis of alkyl halides by aqueous alkalis forms alcohols.

$$R-X+OH \longrightarrow R-OH+X$$

where R may be an alkyl, aryl, metal or metalloid group, and X may be an inorganic or organic anion. The reaction may be

- 1. S_N^2 (substitution, nucleophilic, bimolecular), and
- 2. S_{N^1} (substitution, nucleophilic, unimolecular).

In S_{N²} reactions, the rate of reaction is of second order (involve two particles) or in general, such reactions involve a Walden inversion and there is a 5-co-ordinate intermediate in which both X and say, OH are bonded to same carbon atom.

In S_N^1 reactions the rate of a nucleophilic substitution reaction depends upon the concentration of alkyl halide only, *i.e.*, a first order reaction.

$$R-X \longrightarrow R^+ + X^-$$
 (slow reaction)

For example, the hydrolysis of tertiary butyl bromide to form tert-butyl alcohol.

$$\begin{array}{c}
CH_{3} \\
CH_{3} \\
CH_{3}
\end{array}
C-Br \stackrel{Ionization}{\rightleftharpoons} +C + Br^{-}$$

$$\begin{array}{c}
CH_{3} \\
CH_{3}
\end{array}
+H_{2}O$$

$$\begin{array}{c}
CH_{3} \\
CH_{3}
\end{array}
C-OH \stackrel{-H^{+}}{\leftarrow} CH_{3}$$

$$\begin{array}{c}
CH_{3} \\
CH_{3}
\end{array}
C-OH_{2}$$

$$\begin{array}{c}
CH_{3} \\
CH_{3}
\end{array}
C-OH_{2}$$

$$\begin{array}{c}
CH_{3} \\
CH_{3}
\end{array}
C-OH_{2}$$

The energy required for the cleavage of C-Br bond is supplied by the energy released in the solvation of carbonium and bromide ions.

Nucleosides. Crystalline and sparingly soluble glycosides of heterocyclic bases, mainly of purine and pyrimidines.

Nucleotides. A term used for phosphates of glycosides of all heterocyclic bases to encompass such compounds as adenosine triphosphate, nicotinamide adenine dinucleotide and other coenzymes.

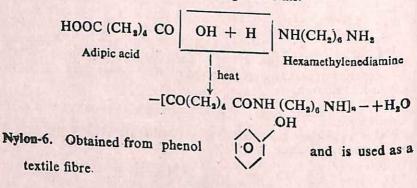
Nucleus. The compact positively charged centre of an atom consisting of one or more nucleons (protons and neutrons) around which is a cloud of electrons. Hydrogen has the simplest nucleus of one proton only.

Nuclide. A term used in radioactive decay.

Nujol. A brand name for a heavy medicinal liquid paraffin,

Nylon. A type of synthetic polymers linked by amide groups
—NH.CH—. Manufactured by condensation of α, ω-aminomonocarboxylic acids or of aliphatic diamines with aliphatic
dicarboxylic acids. Important nylons are:

Nylon-6, 6, [CO(CH₂)₄ CONH(CH₂)₆ NH]_n. Obtained from heramethylene diamine and adipic acid used in monofilaments, textile cables, insulation and in packing materials.





O. Oxygen.

Occlusion. A term used to denote either absorption of a gas by a solid (e.g., the occlusion of hydrogen by Pd) or the absorption of an electrolyte by a precipitate.

Ochre. A variety of the earthy iron-containing pigments with yellow to brownish-red in colour.

Ocimene, C₁₀H₁₆. An alicyclic monoterpene of the structural

Octadecanoic Acid (stearic acid), CH₈ (CH₂)₆ COOH. A solid carboxylic acid present in fats and oils as the glyceride.

Octadecenoic Acid (oleic acid), Cis-9-octadecenoic acid,

 $CH_3(CH_2)_7$ $CH=CH(CA_2)_7$ COOH

A carboxylic acid occurring naturally in fats and oils (as glycerides) and in cow's milk. Used in crude form as resins, lubricant, etc.

- Octabeeral Co-ordination. Co-ordination of six ligands which are present at the coiners of an octahedron, e.g., SF6 has a octahedral co-ordination about S.
- Octane Number or Octane Rating. A rating for the performance of petrol in internal combustion engines. The octane number of a gasoline is 'the percentage of iso-octane present in a mixture of iso-octane and n-heptane, when the mixture has the same knocking performance in the experimental engine as the gasoline under investigation. For example, 2, 2 4trimethylheptane (iso-octane) is given the value of 100 and heptane is given a value 0. So, octane number measures the freedom from 'knocking.' Used chiefy, for motor or aviation
- Octant Rule. A rule, applied specially to steroids which correlates the sign and amplitude of Cotton curves with the geometry and substitution of cyclohexenone derivatives.
- Octet. A group of eight valence electrons (no d electrons). The completion of octet leads to the stability and which is a basis of Lewis Octet Theory. So,
 - 1. the noble gases have complete octets (except He) and
 - 2. the ions of electropositive or electronegative elements are with a complex octet, e.g.

the covalent bonds are formed by sharing to complete

Exceptions: In BF₃, B has 6 electrons and in SF₆, S has 12

Oenanthic Acid, n-Heptanoic Acid, C7H14O2. An oily liquid, B.P. 115°C and present in some natural oil and waxes.

Oestradiol, Estradiol, C₁₈H₂₄O₂. M.P. 174°C.

Oestrogen, Estrogens. The follicular female sex hormone.

Oestrone, Estrone, C₁₈H₂₂O₂. A female sex hormone.

- Ohm (R). A unit of electrical resistance. It is the resistance offered by a conductor when a current of one ampere flows through a potential difference of one volt. 1%=IVA-1 (Ohm's
- Oil. Naturally occurring hydrocarbons which are used as a fuel (after refining) or an oily substance obtained from natural products, e.g., oil of cade.
- Oil of Anise (oil of aniseed). Obtained by steam distillation of anise fruits, Pimpinella anisum or Illicium verum. Used medicinally as an ingradient of cough mixture and as a flavouring
- Oil of Cade (juniper tar oil). Brownish black oil obtained by destructive distillation of the branches of Juniperus oxycedrus. Used medicinally in ointments for skin diseases.
- Oil of Camphor. Two fractions of the natural oil of comphor are
 - (1) Light oil of camphor. A colourless fraction having a small
 - (2) Dark oil of camphor. A fraction containing a large quan-
- Oil of Caraway. Obtained by distillation of the ripe fruits of Carum carvi. Used medicinally to remove flatulence in babies.
- Oil of Cloves. The volatile oil obtained by distillation of cloves which contain a large quantity of equgenol. A carminative and
- Oil of Coriander. An aromatic, stimulative and carminative oil prepared by the distillation of ripe fruits of Coriandrum sativum. Contains mainly coriandrol C₁₀H₁₇OH (65%-80%).
- Oil of Peppermint. An oil obtained by distillation of fresh flowering tops of Mentha piperita which contains about 50% menthol plus menthyl esters plus menthone. Used as carminative,
- Oil of Sandalwood. A volatile oil obtained by the distillation of Santalum album tree. Oil of sandalwood contains about 93%
- Oil of Wintergreen. See methyl salicylate.
- Olefin Complexes. Compounds of olefins and metals (coordinate compounds). Formed by π-bonding orbitals of olefin plus
- Olefins. See alkenes.

Oleic Acid. See octadecenoic acid.

Oleum. See sulphuric acid.

Oligomer. A polymer made up of small number of monomeric units (oligomerization).

Olivine, 9Mg₂SiO₄, Fe₂SiO₄. A silicate having discrete SiO₄²⁻ anions. Used in the manufacture of refractory bricks.

'Onium Compounds. A type of compounds containing R_zA⁺ type of cations (e g., R₄P⁺, R₃O⁺. R₂S⁺, etc.). Analogous to NH₄⁺ ions.

Opal. An almost non-crystalline form of quartz or hydrated silica (up to 12% combined water). Used as a gem stone.

Opium. The dried latex exuded from the unripe capsule of *Papaver Somniferum* after the outer skin had been cut. It is a mixture of alkaloids mainly morphine, codeine and narcotine. Used chiefly as a narcotic.

Oppenauer Oxidation. The oxidation of s-alcohols to ketones in the presence of alumunium tert-butoxide.

Optical Activity. A property of certain substances to rotate the plane of polarization of polarized light. It can be exhibited by some solids, liquids, gases and solutions. It is due to the presence of asymmetry either in a molecule, an ion or a crystal lattice. The asymmetry in a molecule arises due to four different groups joined to one carbon atom, or due to restricted totation about a central bond. These substances are optically active. Optically, active substances are of two types,:

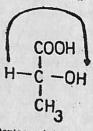
(i) dextrorotatory (d, +) which rotate the plane of polarized light to the right with respect to the direction of the incidence light (clockwise).

(ii) laevorotatory (l, -) which rotate the plane of polarized light to the left with respect to the direction of incidence light (anti-clockwise). The prefix (±) - is used for racemic substances which do not rotate the plane of polarized light. These signs are arbitrary as the rotation can change with wavelength (λ) of light (cotton effect).

In carbohydrates and amino-acids the prefix D- and L- are used to indicate configuration and not the direction of rotation.

For amino-acids the same nomenclature has been adopted. All naturally occurring α-amino-acids have a configuration with -NH₂ group on the left side and by analogy to L-glyceric aldehyde belong to L-family irrespective of their direction in which they rotate the plane of polarized light.

Isomers of lactic acid



Dextro-rotatory (d) or (+) lactic acid

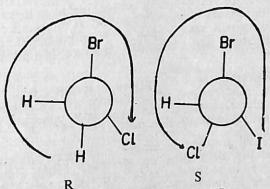
Laevo-rotatory (l) or (—)

R, S Convention for Configuration (Rectus and Sinister system). The two steps involved are:

Step I. Following a set of sequence rules, a sequence of priority is attached to the four different groups of atoms attached to the asymmetric carbon atom. The criterion of priority is based on atomic number—an atom with higher atomic number gets higher priority, e.g., in the case domethane given below. of bromochlerois-

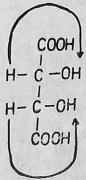
The sequence of priority is I, Br, Cl, H.

Step II. The molecule is next visualised to be so oriented that the group of lowest priority is directed away from us, and the arrangement of the remaining groups observed. Proceeding from the group of highest priority to the group of second priority and hence to the third, our eve travels in a clock wise direction, the configuration is termed R. When eye travels in anti-clockwise direction, the configuration is termed S.



Racemic modification is denoted by prefix RS, e.g., racemic lactic acid is written (RS)-lactic acid.

One structure of tartaric acid has two parts of the molecule which are mirror image of each other (having a plane of symmetry) and is the *meso* form of tartaric acid. This form is not optically active.



Meso-tartaric acid or (dl)-tartaric acid

The optical rotations are measured as 'specific rotations', which are given by $[a]p^{to} = \frac{aobs}{1 \times c}$, where $[a]p^{to} =$ the specific rotation at 1°C, abs=the observed angle of rotation, l=the length of solution in decimeters and C=the concentration (gm/litre). Optical Electrons. The outermost electrons of an atom.

Optical Purity. For an optically active substance, it is the ratio of specific rotation, $[\alpha]_D$ to the specific rotation of pure enantio-

Optical Rotatory Dispersion (ORD). Describing the changes in optical rotation of an optically active molecule (organic or inorganic) with the wavelength of light (λ) (Cotton effect). Plots of radiation versus wavelength (λ) can he used for providing information about the molecular structure of optically active

Optoacoustic Spectrometry. A spectrophotometric technique used to examine most regions of electromagnetic spectrum. When modulated electromagnetic radiation falls upon the sample it produces heat when absorption occurs.

Orbital. A region around an atomic nucleus in which there is a high probability of finding an electron. More accurately an allowed energy level for electrons. Orbital can also be visualized as a cloud of electron charge. The shapes and energies of atomic and molecular orbitals can be calculated by quantum theory. The shapes of atomic orbitals depend on the angular momentum (the sub-shell). For each shell there is:

s-orbital-1 (spherical)

p-orbital-3 (each have two lobes)

d orbital-5 (complex shapes with four lobes)

f-orbital-7

s-orbital can accommodate 2 electrons, p.6 electrons, d-10 electrons and f-14 electrons respectively.

Molecular orbitals are formed by overlap of atomic orbitals.
When the orbital is completely symmetrical about an axis between nuclei, it is a σ orbital, e.g., s-s overlapping in hydrogent molecule or pp overlapping with their lobes along the axis. When the overlapped atomic orbitals fall in regions above and below the axis or overlap at right angles to the axis they for me or is. $pi(\pi)$ orbital, e.g., pp overlapping at right angles to the axis. A double bond has one σ and one π bond.

Hybrid orbitals are atomic orbitals formed by combination of p, d and f atomic orbitals. Carbon combination of s, p, d and f atomic orbitals. Carbon with electronic configuration $1s^2$, $2s^2$ p^2 has one filled s orbital, two filled p orbitals, and one empty p orbital. These four orbitals as one 'empty' p orbital. These four orbitals may hybridize as-

sp³ hybridization. Four equal orbitals arranged tetrahedrally, each with one electron, e.g., in methane,

sp2 hybridization. One s and two of, the p orbitals hybridize and act as three orbitals in a plane at 120°. The remaining p orbital is at right angles to the plane and can form π bond.

sp hybridization. Forming two orbitals in a line.

Bonding orbital. Combination of two atomic orbitals which has a concentration of electron density between the nuclei and are responsible for holding the nuclei together.

Anti-bonding orbital. Combination of two atomic orbitals that has slightly higher energy and tends to repel atoms.

- Orbital Angular Momentum. The angular momentum of a set of electrons is an atom or ion (L=0, for empty and completely filled shell).
- Orbital Moment. The component of the magnetic moment of an atom or ion due to the motion of an electron around nucleus.
- ORD. Optical rotatory dispersion.
- The sum of the powers to which the concen-Order of Reaction. tration terms are raised in the rate equation of a reaction at a specified temperature. For a reaction,

$$aA+bB\rightarrow Products$$

the rate equation is

$$\frac{\mathrm{d}x}{\mathrm{d}t} = k [A]^a [B]^b$$

and the order of the reaction is (a+b). Here, a and b denote the number of moles A and B. Order of a reaction is an experimentally determined quantity and depends upon the mechanism of reaction, temperature of reaction and so on. For example, thermal decomposition of N₂O₅ is a first order reaction, though it seems to be a second order reaction.

$$2N_2O_5 = 2NO_2 + O_2$$

This reaction proceeds in two steps:

(i)
$$N_2O_5 = 2NO_2 + O \text{ (slow)}$$

$$(ii) O+O=O_2 \qquad (fast)$$

The slowest step in a multistep reaction is the rate determining step.

Ore. A mineral source of a chemical element.

Ore Dressing. A method for concentrating the important part of the ore from the original mineral by physical means.

Organic Chemistry. The chemistry of carbon compounds. Originally the term 'organic chemistry' was applied to substances produced by or otherwise associated with living matter from animal or vegetable origin. Nowadays, the organic chemistry is the study of the compounds of carbon, whether they are isolated from natural sources or synthesized in laboratory. Most organic compounds contains C and H as essential elements and other elements like N, S, P, halogens, etc., as extra elements. Organic compounds generally contain covalent bonds and differ from inorganic compounds which contain typical ionic links. Metallic carbonates and some other compounds are inorganic.

Organoboranes. Compounds of carbon and boron having at least one direct C to boron bond, e.g., trialkylboranes (R3B) and dialkylborane (R,BH).

Organoborates. Compounds of boron containing BR₄ anions. Some organoborates may be polymeric, e.g., LiBMe₄.

Organo Chlorine Insecticides. Highly active insecticides.

Organolithics. Constituted by organics and silicates.

Organometallic Compounds. Compounds containing a carbon metal bond, e.g., tetraethyl lead, (C2H5)4Pb, Grignard's reagent,

Organometalloids. Compounds having a direct bond between one or more carbon atoms and one or more metalloid atoms.

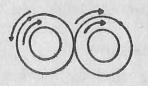
Organophosphorus Compounds Compounds having C-P bonds and which are used as pesticides, e.g., malathion, parathion and dimefox. Some esters and thio-ethers also belong to this class of compounds.

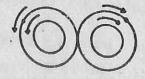
Organosol. Colloidal particles dispersed in organic medium. For example, alcohol, benzosol (benzene medium), etc.

Orgel Diagrams. Graphs denoting the relation between the energies of various electronic states and the crystal field splitting.

Orsat Gas-Analysis Apparatus. An apparatus used in gas analysis.

Ortho hydrogen. When the nuclei of a hydrogen molecule spin in the same direction, para hydrogen is formed, but when the spins are in opposite direction ortho hydrogen is formed. Ordinary hydrogen is a mixture of two forms (25% para and 75% ortho) at ordinary temperature. Ortho hydrogen is prepared by cooling ordinary hydrogen to a very low temperature in con-





ortho-hydrogen

para-hydrogen

Para hydrogen always tends to revert in the ortho-form (morestable form). The process is accelerated

- (i) By heating 800°C or more.
- (ii) By treatment with catalyst like Pt or Pd.
- (iii) By passing an electric discharge.

The o-and p-hydrogen differ in their physical properties but show identical chemical properties.

Orthophosphoric Acid. See Phosphoric (V) acid.

Orthophosphorus Acid. See Phosphoric acid.

Orthorhombic System. A crystal system in which a unit cell hasthree unequal axes at right angles.

Os. Osmium.

Osazones. Organic compounds with the group

They are produced by reacting a-diketones, a-hydroxyaldehydes, hydroxyketones, aminoaldehydes or aminoketones with arylhydrazine such as phenylhydrazine. Sugars form osazone derivatives with sharp melting points (a test for sugars), e.g., glucose (C₆H₁₂O₆) forms glucosazone

(CH₂OH. (CHOH)₅ C: N.NH. C_6H_5 . CH: N.NH. C_6H_6).

- Osmic Acid. The name of osmium tetraoxide and its solution in water. Free OsO4 is present in solution.
- Osmiridium. Native or synthetic alloys of osmium (15-40%) and iridium (50-80%). Used for making sparking agent and nibs
- Osmium, Os. At. No. 76, At. Wt. 190.2 M.P. 3045°C, B.P. 5027°C, mium, Os. At. No. 76, At. wt. 190.2 M.F. 5043 C, B.P. 5027°C, D. 22.6. A transition metal of platinum group. Occurs as a sulphide or as osmiridium. The electronic configuration is 5d² sulphide or as osmiridium. States from +8 to 0. Forms OsO₄ with 6s². Shows oxidation states from +8 to 0. Forms OsO₄ with O₂ at high temperatures. Used for hardening alloys of plati-
- Osmium Oxides. OsO4 is a colourless toxic material obtained from Os and O_2 at high temperature. M.P. 25°C. Os O_2 is a dark coloured oxide obtained from Os plus OsO4.
- Osmosis. The phenomenon of diffusion of a solvent through a semipermeable membrane from the solvent to a solution or from a dilute solution to a concentrated one is called osmosis.

Osmotic Pressure (n). The osmotic pressure of a solution is the pressure required to prevent osmosis when the solution is separated from pure solvent by a semipermeable membrane. It is a colligative property and the magnitude of the osmotic pressure depends on various factors such as temperature, concentration, etc. but is independent of the nature of the membrane. When the equilbrium is established between the two sides of the membrane, the rate of diffusion of solvent in the two opposite pressure (n) is given by

$$\pi = \frac{cRT}{M}$$

where c is the concentration of solute, M is the molecular weight of solute (in grammes per unit volume), R is the gas constant and T is the absolute temperature.

Ostwalds Ripening. A process of crystal growth.

Ovalbumin, Egg Albumin. The main protein constituent of white of egg.

Over-voltage, Over-potential. The difference between the potential of the electrode when gas evolution is actually observed and the theoretical reversible value for the same solution is called overabove that of the reversible electrode, that must be applied to an electrode for electrolysis to proceed at a measurable rate. Overreduction, polarography, etc.

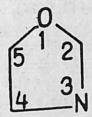
Oxalic Acid, Ethaeudioc Acid, C₂H₂O₄, COOH. Colourless pri-

smatic crystals, soluble in water, poisonous, and crystallizes from water as a dihydrate. M.P. 101°C (hydrous) and 189°C (anhy-Manufactured from sodium methanoate. Decomposed to CO, NH₄+ salts are water soluble. Oxalic acid is used for textile finifrom cotton.

Oxamide $C_2H_4N_2O_2$, $(CONH_2)_2$. Colourless crystals which form cyanogen with P_2O_5 .

Oxanthrol. A tautomer of anthraquinone.

Oxazole Ring. A cyclic ring as shown.



Oxetane Polymers. Polymer derivatives of oxetane.

(CH2CH2CH2O) containing repeating unit (CH2CR2CH2O)n

Oxidases. Dehydrogenating enzymes which transfer hydrogen directly to molecular oxygen.

Oxidant. An oxidizing agent, e.g., K₂Cr₂O₇, KMnO₄, KClO₃, etc. In rocket fuels, the oxidant provides oxygen for combustion, e.g. liquid oxygen or H₂O₂.

Oxidation. A process of removal of electrons from an atom or ion, e.g.,

$$Fe \rightarrow Fe^{2+} + 2e$$

 $Fe^{2+} \rightarrow Fe^{3+} + e$

and the oxidation state is increased. The oxidation state of an atom is indicated by the number of electrons lost or effectively lost by the netural atom i.e., the oxidation number. The oxidation number of a negative ion is negative. Oxidation and reduction go side by side.

β-Oxidation. The process of degradation of fatty acids in biological systems to produce acetyl-CoA which is fed into the citric acid cycle to produce ATP, etc., eventually.

Oxidation State or Oxidation Number. It is the charge which an atom appears to have when the electrons are counted according to the arbitrary rules. The rules are:

- 1. The electron shared between two unlike atoms are counted with the more electronegative atom.
- 2. The electrons shared between two like atoms are divided equally between the sharing atoms.

For example, chromium is $K_2Cr_2O_7$ molecule has oxidation state of +6. The oxidation number of Mn in KMnO₄ molecule is +7.

Elements are in the zero oxidation state.

Oxidative Addition. Describing a class of reactions involving direct addition to an element or multiple bond with increase in oxidation state e.g., addition of Cl₂ to PCl₃ to form PCl₅.

Oxidative Phosphorylation. A process occurring along the respiratory or electron transport chain resulting in the generation of ATP.

Oxide. A compound of an element with oxygen. Oxides are divided into four categories.

- 1. Acidic oxides. They react with bases to form salts. e.g., SO₂, CO₂, P₂O₅, etc.
- 2. Basic oxides. There react with acids to produce salts, e.g., CaO, Ba, CuO, etc.
- 3. Atmospheric oxides. They exhibit both acidic and basic properties, e.g., Al₂O₃.
- 4. Neutral oxides. They do not react either with acids or bases, e.g., NO and CO.
- 5. Peroxides. They are metallic derivatives of H₂O₂. Some oxides contain metal atoms in two or more oxidation states. Mixed oxides are obtained by fusing two or more metal oxides (no discrete oxyanion).

Oxidised Rubber, Rubbon. A type of rubber produced by oxidation of rubber with air in the presence of a catalyst can be vulcanised and chlorinated. Used for impregnation, as an adhesive and for electrical insulation.

Oxidising Agent. Substances which carry out oxidation, e.g. MnO₄ and Cr₂O₇²⁻ in acid oxidizes Fe²⁺ to Fe⁸⁺ or C₂O₄²⁻ to CO₂ and water.

Oximes. Organic compounds having the group=N.OH which is united directly to a carbon atom.

Oxine. 8-Gydroxyquinoline.

Oxirane (epoxide). A name for the heterocycle compound.

Oxonium. A positive ion having a central oxygen atom, e.g., hydroxonium ion (H₃O⁺), trimethyloxonium (Me₃O⁺). Used as alkylating agent for nucleophlies.

Oxo Reaction. A reaction in which the water gas or synthesis gas (H₂+CO) adds to an alkene under pressure in the presence of a catalyst.

Oxyanions. Anions formed by co-ordination of oxide ions, O²⁻, to metal or non-metal cation, e.g. the SO₄²⁻ ions (S⁶⁺+4O₂⁻).

Oxyazo Dyes. Azo dyestuffs containing a hydroxyl group.

Oxycyanogen, (OCN)₂. A pseudohalogen (KOCN+Cl₂).

- Oxygen, O. At. No. 8, At. wt. 15'9994, M.P.-218'4°C, B.P. -182'162°C. A colourless diatomic gas, the first member of group VI of the periodic table. Electronic configuration 1s2, 2s2, 2p4. Forms dinegative ion O2 or two covalent bonds. Occurs in air (21%) by volume and in lithosphere 47% by weight. Manufactured by distillation of liquid air and prepared by heating KClO₃ with MnO₂. Forms oxides such as, K₂O, BaO etc. Exists as paramagnetic O2 and as ozone, O3. Forms compounds with zero group elements, e.g. XeO₃, XeO₄, XeOF₂, XeO22-. Also forms several type of oxides, e.g. acid oxide, basic oxide, etc. Oxygen occurs in three natural isotopic forms, 16O, 17O and 18O. Its oxidation state in oxides is -2.
 - Oxygen Carrier. A molecule which can react directly with oxygen (molecular) and subsequently use this oxygen for oxidation of a species, e.g. in the haemoglobin.
 - Oxygen Cathode. A Pt or Au electrode for measuring oxygen content by measuring the rate of cathodic reduction of O₂. This electrode is used in amperometry.
 - Oxygen Fluoride, OF₂. Obtained from F₂ and dilute cold NaOH. Used as a strong oxidizing agent when sparked.
 - Oxythallation. Thallium (III) salts used as oxidizing agents.
 - Oxytocias. A cyclic peptide hormone which stimulates contraction of the smooth muscle of the utreus. It is secreted by the posterior lobe of the pituitary gland.
 - Ozokerite. A natural mineral wax which forms ceresin wax (hard
 - Ozone (trioxygen, O₃). An allotrope of oxygen prepared by passing O₂ through a silent electric discharge, M.P.—193°C, B.P.-1120°C. Present in upper atmosphere and protects earth's surface from ultra-violet radiation. Ozone is a strong oxidizing agent. Forms ozonides with alkenes. Used for oxidation, sterilization and
 - Ozonides. The addition of ozone to various types of unsaturated organic compounds produce ozonides. Prepared by passing ozonized air or O₂ into the unsaturated compounds dissolved in a suitable solvent. The structure is as follows:

Ozonides are explosive, thick oils and readily decomposed by water ethanoic acid, etc. Used to detect the position of double bond in unsaturated compound.

Ozonizer. An apparatus for the preparation of O3.

Ozonolysis. The addition of O₃ to an unsaturated organic compound to form ozonide which is unstable and is decomposed in situ.

P

P. Phosphorus.

Pa. Protactinium.

Packed coloumn or Packed Tower. Used for gas absorption, for liquid-liquid extraction, for industrial distillations, etc.

Packing Fraction. Aston found that in most cases there was a slight deviation from whole number for the mass of an isotope. The nearest integral number called the mass number (A) is integral to the numbers of protons and neutrons. The deviation from whole number varies from nucleus to defined as

packing fraction = isotope mass - mass number ×10⁻⁴

Negative fraction implies stability of nucleus while positive fraction indicates unstability of nucleus. For example, the chlorine isotope of mass 35 has the packing fraction,

$$\frac{34.980 - 35.000}{35.00} \times 10^4 = -5.7$$

For the lightest and heaviest elements, the packing fraction is fraction. The packing fraction is fraction.

Paint. A covering material applied to wood, metal etc., to improve apperance and also to protect the substrate against rust, etc.

Palladium, Pd. At No. 46, At. wt. 106.42, M.P. 1552°C, B.P. 3140°C, D 12.02. A transition metal occurring in platinum ores.

An element of the nickel group, electronic configuration 4d10. The metal is dissolved by Conc. HNO3 or hot H2SO4. Forms organometallic derivatives which are less stable than for Pt. Used in alloys, e.g. white gold, as a catalyst for hydrogenation. etc.

- Palladium Ammines. Pd (II) forms stable amino-salts which resemble bivalent platinum salts and have the co-ordination number 4. Represented by the formula (Pd A₄)X₂ and [Pd A₂X₂] where A is ammonia or a substituted ammonia and X a univalent acidic radical.
- Palladium Black. A very finely divided precipitate obtained by the action of reducing agents on palladium solutions.
- Palladium Halides. Palladium forms halides with halogens such as, a brick-red PdF4, black PdF3, violet PdF2, PdCl2 (Pd plus Cl2), PbBr2 and PdI2.
- Palladium Hydrides. Pd absorbs hydrogen to formhydrido complex e.g. [Pd HBr (PEt3)2].
- Palladium Oxide PdO. The only oxide, obtained from Pd plus O, at 800°C.

Palmitic Acid. Hexadecanoic acid.

Palygorskite. A type of fuller's earth.

- Pamoic Acid, Embonic Acid, 2.2' dihydroxy-1,1'-dinaphthyl-methane-3,3"-dicarboxylic acid, C23H16O6. Used in medicines.
- Panchromatic Sensitization. To make photographic sliver halides emulsion more sensitive to the whole of the visible spectrum by
- Pantothenic Acid, CoH17NO5. A vitamin, oil and a constituent of
- Papavevine, C₂₀H₂₁O₄N. One of the alkaloids of opium. Insoluble in water, soluble in hot alcohol, optically inactive. Causes tetanus if taken in excess.
- Paper. Dehydration product of an interwoven mat of hydrated cellulose fibres. The properties of papers depend upon the degree of hydration and sub-division of fibres.
- Paper Chromatography. A technique used for the analysis. A special paper acts as a stationary phase and a suitable solvent as special paper acts as a stationary plane is spotted on to the a mobile phase. A small sample of mixture is spotted on to the a mobile phase. A small sample vertically in mobile phase. The paper is placed vertically in dissolves in this paper. The paper is placed dissolves in this mobile components within the sample mixture dissolves in this mobile phase and are carried up by the paper. The paper is removed phase and are carried up of and dried. The paper is developed to locate the position of

colourless fractions by spraying with a suitable chemical, e.g., ninhydrin or by explosure to u.v. radiation. It is an application of partition law.

Para (p-). A pretix used to name disubstituted benzene derivatives. A paraposition is at carbon no. 4 with respect to carbon number 1 in a benzene ring.



Parabens. Alkylhydroxybenzoates used as food preservatives.

Paracetamol, 4-Acetamidophenol, C₈H₉NO₂. A white powder analgesic and antipyretic. M.P 169-172°C.

Parachor, P. An additive property defined as

$$P = \gamma^{\frac{1}{2}} \left(\frac{M}{D - d} \right)$$

where γ is the surface tension, M is the molecular weight, and D and d are the densities of liquids and vapours respectively. Of compounds.

Paracyanogen, (CN)x. A white or brown polymer obtained by heating cyanogen to 400°C.

Paraffins. See alkanes.

Paraffin Wax. See petroleum wax.

Paraformaldehyde, Paraform, Trioxymethylene. A mixture of polymethylene glycols of the type $(CH_2O)_nH_2O$ where n is 6 to 50. A white amorphous powder with the odour of formaldehyde. Forms methanol on heating. The commercial product contains 95% methanol.

Para-hydrogen. See ortho-hydrogen.

Paraldehyde (C₂H₄O)₃. A colourless, mobile liquid, slightly soluble in water with M.P. 125°C and B.P. 124°C.

Paramagnetism. A property exhibited by an ion or molecule which contains unpaired electrons.

Paraquat, 1-1'-dimethyl-4.4'-bipyridylium dimethyl sulphate or dichloride. A contact herbicide, M.P. 175-180°C. Poisonous by ingestion and causes irreversible lung damage.

Parathion, C₁₀H₁₄NO₆PS. An insecticide.

Paris Green (Schweinfurter-green). A brilliant pigment with the compositition $Cu(C_2H_3O_2)_2$, $3Cu(AsO_2)_2$. Used as an insecticide.

Parkerizing. The phosphate treatment of steel.

Partial Pressure. In a mixture of gases or vapours it is the pressure which the same amount of the constituent would exert if it were present alone in a vessel of the same volume as that occupied by the gaseous or vapour mixture. In a mixture of two gases A and B

aseous of April and
$$p_B = \frac{n_B}{n_A + n_B}$$
. P and $p_B = \frac{n_B}{n_A + n_B}$. P

where pA and pB are the partial pressures of A and B respectively, n_A and n_B are the number of moles of A and B in the mixture and P is the final pressure.

Parting. A process used to separate Au from Ag either by dissolving Ag in Conc. HNO₃ or H₂SO₄ or the perferential reaction of Ag with Cl2 to form AgCl.

Partition Column Chromatography. See Chromatography.

Partition Co-efficient. When a solute dissolves in two non-miscible liquids, the partition co-efficient is the ratio of the concentration of one liquid to the concentration of the other liquid and it is or one liquid to the concentration of concentration the activities of the solute in two solutions are used.

If C_1 and C_2 represent the concentrations of a solute in the **two** immisible liquids, then $\frac{C_1}{C_2} = K$

where K is the Distribution Co-efficient or partition co-efficient of the solute between the two liquids at the given temperature.

PAS. 4-Aminosalicylic acid.

Paschen Series. A series of lines in the 1R spectrum emitted by hydrogen atoms. These lines correspond to the radiation emitted when electron falls to third energy level (lowest). The wavelength (λ) of the series is given by

$$\frac{1}{\lambda} = \mathbb{R}\left(\frac{1}{3^2} - \frac{1}{n^2}\right)$$

where n is an integar and R is the Rydberg constant.

Passive or Passivity. A process which renders the metal surfaces unreactive due to the formation of surface layer oxide (oxides) incontact with strong oxidizing agents, For example, Fe, Co, Ni etc., in contact with acids such as HNO₃ or chromic acid become passive due to the formation of a thin oxide layer.

Pasteur Effect. The decomposition of sugar by yeast and other cells is greater in the absence of O₂(CO₂+C₂H₅OH) rather than its presence (CO₂+H₂O). This is known as Pasteur effect.

Pasteurization. It is the process for removing some non-sporeproducing organisms from milk, butter, beer at low temperatures (63°C for half an hour and 72°C for 20 seconds).

Patina. The formation of an oxide film on the surface of e.g., bronze, iron, etc., resulting in a decorative and corrosion-resistant surface.

Pauli Exclusion Principle. In an atom no two electrons can have number is always $\pm \frac{1}{2}$).

Pb. Lead.

PCB. Polychlorinated biphenyls.

Pd. Palladium

Pearl Ash. See potassium carbonate.

Peat. A brown, fibrous mass of partially decayed plant matter.

It occurs in nature and is used for firing in power stations.

Converted to peat charcoal by low temperature carbonization.

Pectins. A group of polyuronides presents in fruits of all types chiefly in citrous fruits. It is a mixture, having methyl pectate

as the chief constituent. Manufactured as a white powder, soluble in water and is used to assist the setting of jams and iellies.

Pelargonic Acid, Nonanoic, Acid, C9H18O2, CH3(CH2)7. COOH. An oily liquid found with other fatty acids and obtained from oleic acid. M.P. 12.5°C, B.P. 253-254°C. Also present in human hair.

Penicillin. A group of bactericidal antibiotics having the basic ring structure of penicillanic acid.

Penicillinase. An enzyme present in a bacteria which hydrolyses the β-lactam ring of penicillin to penicilloic acid.

Pentaborane, B5H9; B5H11.

Pentachlorophenol, C₆Cl₅OH. Prepared by chlorinating phenol. Used as a food preservative and as a fungicide

Pentnerythritol tetranitrate, PETN, Penthrite, C(CH₂ O NO₂)₄. M.P. 141°C. A powerful explosive. Forms pentolite with TNT. Used as a pain reliever for angina pectoris.

Pentanes, C₅H₁₂. Three isomeric forms of the compound. Occur in the low boiling fraction of petroleum. Inflammable and used as a standard illuminant in photometry.

Pentanoic Acid (Valeric acid), CH3(CH2)3COOH. A colourless liquid carboxblic acid. Used in the preparation of per fumes.

Pentavalent. Describing a valency of five.

Pentose (aldopentose). A carbohydrate with 5 carbon atoms. For example, D-form of xylose or ketopentoses, etc.

Pepsin, Pepsinogen. A protein digesting enzymes of the gastric juice which is secreted by gastric cells of vertebrates.

Peptidases. Proteolytic enzyme responsible for breaking peptide

Peptide. Substances made up of two or more amino-acids which are linked by the peptide bond —CO.NH—. They are di-tri-, are linked by the peptide bolls to the number of amino-acids oligo-, poly-peptides according to the number of amino-acids linked by the peptide bond. Glycine is the simplest peptide (H2N.CH2 CO.NH.CH2.COOH).

Peptization. A process of converting a fresh precipitate into a colloidal state by adding an electrolyte. The electrolyte added is called a peptizing or dispersing agent. For example, a reddish ferric hydroxide sol is obtained by adding a small quantity of FeCl₃ solution to freshly precipitated Fe(OH)₃ and stirring.

Fe $(OH)_3+Fe^{3+} \rightarrow Fe(OH)_3$: Fe^{3+} Precipitate Sol

In general, peptization is a process of converting a solid state into its stable colloid by adding suitable agents. It can also be carried out by continuous washing of precipitate with water; e.g. Formation of BaSO₄ colloid by continuous washing with water.

- Peptenes. Proteins are degraded (e.g. by enzymes) to form low molecular weight proteins which are known as peptones.
- Per-acids. Per oxy-acids are obtained by replacing oxide in oxy-anions by $(O-O)^{2-}$ or $(O-OH)^-$ groups. Prepared by sulphuric acid $(H_2S_2O_5)$ or per-chromic acid $(H_2O_2$ plus $K_2Cr_2O_7$ way, e.g. per-benzoic-acid.
- Perborates, Perborax. Sodium perborates used as a bleaching agent and antiseptic.
- Percarbonates. Carbonates having varying amounts of H₂O₂ of crystallization, e.g. 2Na₂CO₃, 3H₂O₂, Used in laundering for cleaning dentures.
- Perchlororates, MClO₄. Obtained by electrolytic oxidation of (ClO₃)⁻ and contain (ClO₄)⁻ ion. For example, KClO₄.
- Perchloric Acid, HClO₄. A colourless or yellowish liquid obtained from KClO₄ plus H₂SO₄. A strong oxidizing agent which fumes in air. Forms ionized salts.
- Perchloroethylene (tetrachloroethane), Cl₂C=CCl₂. Used as a
- Perchleryl Fluoride, FClO₃. A chlorine oxide fluoride.
- Perfect Gas. A gas that obeys gas laws under all conditions of temperature and pressure.
- Perfect Solution. A solution in which the molecules of the dissolved substance have either no mutual interaction or with the molecules of the solvent.
- Perfluoroalkyl Derivatives. Organic derivatives obtained by replacing all hydrogen atoms, except hydrogen atoms in the functional

- group from the compound. For example, CCl₃COOH (trichloroacetic acid) and C₆H₅COOH (perfluorobenzoic-acid). Used as surfactants (sulphonic acids) and in drug chemistry.
- Perfumes. Natural and synthetic materials such as essential oils, gums, flower oils, etc. and fixatives which make the perfume long lasting.
- Peri. Denotes 1, 8 position on the naphthalene ring.
- Periclase, MgO. Occurs in nature or prepared by heating Mg(OH)2. Used in furnaces.
- Period. A horizontal row in the periodic table. A period has 8 elements from an alkali metal to a noble gas. The periods are classified as follows:
 - (1) the very short period, H and He;
 - (2) short periods, Li-Ne and Na-Ar;
 - (3) two long periods having transition elements K-Kr and
 - (4) a very long period having the rare earth elements Cs-Rn,
- (5) an unfinished period from Fr onwards.
- Periodic Acids H₅IO₆ periodic acid is the crystalline substance (by low temperature electrolysis of a concentrated solution of iodic acid in a cell) soluble in water, strong oxidising agent and loses water on heating. Probable composition is IO(OH)₂. Partially dehydrated acid H2IO5 is also known.
- Periodic Law. According to Mendeleef, the properties of elements are periodic functions of their atomic weights'. If the elements are arranged in order of increasing atomic weights, elements are arranged in order of interests occur at a fixed intervals. A modisnowing similar properties that the elements are the periodic fication of this law stating that the elements are the periodic functions of their atomic number' forms the basis of the
- Periodic Table. A table of elements arranged in order of atomic numbers to show similarities in chemical behaviour. Horinumbers to show similarities in Sichilari centaviour. Horizontal rows are known as periods (7) and vertical columns of zontal rows are known as personal relations are elements having similar outermost electronic configurations are known as groups (one zero group and 7 other groups). Down a known as groups (one zero group and size and electropositive group there is an increase in atomic size and electropositive group there is an increase in the electronegativity increases from behaviour. Along a period the electronegativity increases from left to right in the periodic table (also metallic nature).
- Peritectic. A type of solid liquid phase equilibrium system in which the two components form limited ranges of solid solution which the two components for a completely miscible together in each other, although they are completely miscible together (binary alloys).

Perkin's Reaction. A reaction in which an aldehyde condenses with the sodium salt of fatty acids or their derivatives. Generally, the condensation takes place at a carbon atom. For example, benzaldehyde condenses with sodium acetate to form sodium

$C_6H_5CHO+H_3C.COONa\rightarrow C_6H_5CH=CHCOONa.$

- Perlite. Originally the name given to a glassy volcanic rock. Nowa-days also implies to similar lightly siliceous volcanic materials containing 2-6% of water. Used as a light weight aggregate in heat and sound insulating plasters, in light-weight concretes,
- Permanganates. Inorganic salts containing [MnO₄] ion. They are purple coloured. For example, KMnO4 (obtained from pyrolusite +KOH+air (at 300°C) followed by treatment with CO2) and NaMnO₄, 3H₂O (used as a Condy's fluid in solution form).
- Permanganate Titrations. KMnO₄ is an important oxidizing agent (in presence of mineral acids) used in volumetric estimation of reducing agents. Such as, Fe²⁺ C₂O₄²⁻, H₂O₂, etc, It is a self indicator in most of the volumetric titrations. An acidic solution Mn is reduced M2+ state. It is also used for estimations in neutral solutions, eg., determination of Mn(II) (the Volhard method); the volumetric titration is carried out in the presence

Permanganic Acid. HMnO4.

- Permonosulphuric Acid, Caro's Acid, H₂SO₅, HOS(O)₂OOH. A crystalline acid when pure and obtained by reacting conc. H₂SO₄ and H2O2 or by hydrolysis of H2S2O6 with H2SO4. A powerful
- Permutit. A zeolite of a complex compound, sodium aluminium silicate used for softening hard water by replacing Ca2+ and Mg2+ by Na+ ions. Zeolite is recovered back with NaCl solution.
- Perovskite. Mineral CaTiO3.
- Perovskite Structure. Describing a structure adopted by many ABX3. compounds, where A and B are positive ions and X is F or O2- (negative ions). The structure has A at the corner of a cube, B at the body centre and X at the face. A has 12X neighbours and B has 6X neighbours, e.g., BaTiO3.
- Peroxidase. The enzymes present in plants which are used to
- Peroxides. An oxide containing the O-O2- ion and also a compound containing the -O-O-group, e.g., Na₂O₂, H₂O₂, etc.
- Persulphuric Acid. Caro's acid.

- Peroxides, Organic. Organic substances used as curing agents for resins and elastomers, as a source of free radicals for free radical polymerization, etc., such as dibenzoyl peroxide (acidchloride or anhydride plus Na₂O₂ or H₂O₂).
- Peroxychromium Compounds. CrO(O₂) [H₂O₂+(CrO₄)²⁻] a blue chromium compound extracted into ether.
- Perspex. Trade name for cast polymethyl methacrylate sheet (an acrylic resin).
- Perseitol, C7H16O7. A helptol, M.P. 188°C with (a)p-1'3°.
- Persulphuric Acid, Perdisulphuric Acid, H2S2O8, HO.S(O)2OOS(O)2OH. A di basic acid obtained as salts by electrolysis of sulphates at low temperatures and high current density. The solid persulphates are quite stable and strong oxidizing agents. The free acid is prepared in solid form from chlorosulphonic acid and Caro's acid.

Perxenates . Xenates (VIII).

Perylene. A polycyclic aromatic hydrocarbon.

Peta, (P). A prefix denoting 1015.

Pesticides. See insecticides.

- Petalite, (LiNa) (AlSi₄)O₁₀. Aluminosilicates (2-4.5% Li₂O) used as a source of Li.
- Petrochemicals. Chemicals obtained from crude oil or natural gasor refinery gases or wax, etc.
- Petrol. Motor fuel, gasoline or aviation fuel, etc.
- Petrolatum. A residue from non-asphaltic petroleum, e.g., petroleum jelly, mineral jelly, soft paraffin, etc.
- Petroleum. A mixture of hydrocarbons (or oil products) formed originally from marine animals or plants and found beneath the ground trapped between the layers of rock.
- Petroleum Coke. A hard coke, free from ash and is produced as a by-product from cracking of petroleum oils and also from the distillation (without steam) of heavy wax residues.
- Petroleum Ether, Light Petroleum. Lower aliphatic hydrocarbons volatility distillation range e.g. 40-60°C and 60-80°C ranges (C₅ and C₆ hydrocarbons respec-
- Petroleum Wax. Solid and waxy hydrocarbons obtained from crude oils. Two types of wax are:

- 1. Paraffin wax has macrocrystalline structure wax (n-alkanes, C₂₀H₄₂ and upwards plus small quantities of iso and cycloalkanes).
- Microcrystalline wax (iso-and cycloalkanes plus some aromatics present in lubricating oil residue).

Used in the manufacture of candles, polishes, ointments, etc.

Pewter. An alloy of tin containing 75% Sn and 25% Pb.

Pfeiffer Effect. The change in rotation of a solution of an optically active substance on the addition of a racemic mixture of an asymetric compound.

Ph. The phenyl group, C6H5-.

pH. The negative log of hydrogen ion concentration, i.e. $PH = -\log C_{H^+}$, or more accurately = $-\log C_{H^+}$ (a denotes the activity).

Phalloidins. Amanitins.

Phase. A physically distinct but homogeneous part of a system separated from other parts by boundary surface, eg. water in contact with its vapours form two phases (water liquid and water vapours). All gases always constitute a single phase.

Phase Diagram. A diagram showing the conditions of equilibrium between various phases of a system, e.g., P-T diagrams in one component systems, T-X diagrams (at constant P) or P-X diagrams (at constant T) in binary systems.

Phase Rule. Derived by Willard Gibb's (1876) for a chemical system and is given by the equation

$$P+F=C+2$$

where P is the number of phases, F is the number of degrees of freedom and C is the number of components of the system.

Phase-transfer Chemistry. Describing reactions between heterogeneous systems which are carried out by adding a catalytic agent which transfers a reactant across the interface into the system

Phenacetin, N-acetyl-p-phenetidine, C₁₀H₁₃NO₂. White crystals obtained from phenol. M.P. 137-138°C. Used as an antipyretic analgesic. Shows chronic toxicity towards the kidney.

Phenacyl. The trivial name for the group

$$\begin{matrix} O \\ \parallel \\ C_6H_5-C-CH_2-\end{matrix}$$

o-Phenanthroline, 4,5-Phenanthroline,

C₁₂H₈N₂. M.P. 117°C (anhydrous) and 94°C (monohydrate). Used as a complexing reagent in the estimation of Fe²⁺ and in some chelates of metals.

Phenazone, Antipyrin, 1-phenyl-2, 3-dimethyl-5-pyrazolone, C₁₁H₁₂ON₂ White crystals, M.P. 114°C, B.P. 319°C. Soluble in water. Used as an analgesic and antipyretic.

Phenobarbitone, Phenylethylbarbituric Acid, C12H12N2O3.

White crystals (ethyl ester of phenylethylmalonic acid is condensed with urea), M.P. 174°C. A strong hypnotic and is used as a sedative

Phenol, C₆H₅OH (carbolic acid). A white crystalline solid, soluble in water at room temperature and the hydroxyl group (—OH) is bound directly to C atom of the ring. Acidic nature because of the presence of electron withdrawing effect of the aromatic of the presence of electron withdrawing effect of the aromatic ring. Does not show the typical behaviour of alcohols. Prepared by the following methods:

By fusing sodium salt of the sulphonic acid with NaOH:
 C₆H₅SO₂.ONa+2NaOH→C₆H₅ONa+Na₂SO₃+H₂O
 2C₆H₆ONa+H₂SO₄→2C₆H₅OH+Na₂SO₄.

Raschig process. Catalytic vapour phase reaction of steams and chlorobenzene at 500°C.

3. Dow Process. Hydrolysis of C6H5Cl with NaOH at 400°C

$$C_6H_5Cl+2NaOH \xrightarrow{-H_2O,-NaCl} C_6H_5O+Na \xrightarrow{HCl} C_6H_5OH+NaCl.$$

Phenol is acidic and forms metallic salts. It can be easily halogenated, sulphonated and nitrated. Used in the formation of formaldehyde plastics, e.g. Bakelite, for making nylon (via cyclohexenol), for making picric acid, phenolphthaleine and other dyes and Dettol (a powerful antiseptic).

Phenol Aldehydes, ArCHO. Aromatic aldehydes used in perfumery, electroplating, flavouring, etc.

Phenol Formaldehyde Resins. Resins of phenols (or other aromatic hydroxyl compounds) with methanal (CH₃CHO). Used for moulding coatings, etc.

Phenolic Resins. A type of polymers obtained from phenols and aldehydes.

Phenolphthalein, C₂₀H₁₄O₄. Colourless crystals, M.P. 254°C. An indicator with pH range 8'3 (colourless) to 10'4 (red). Used in medicines as purgative.

Phenoi Red, Phenoisulphonaphthalein, C₁₈H₁₄O₅S. An indicator with pH range 6.7 (yellow) to 8.3 (red). Prepared by heating phenoi functions. Also used as a test for renal

Phenols. Organic aromatic compounds containing at least one —OH group attached directly to the benzene ring. Polyhydric alcohals are also known, e.g. catechol [C₆H₄(OH)_c].

Phenothiazine, C₁₂H₂NS. A yellow crystalline which darkens an cines. Also used to prepare tranquillizers.

Phenoxy Resins, Polyhydroxyethers. Linear thermoplastics used for containers.

$$-(C_6H_4C(Me)_2C_6H_4OCH_2CH(OH)CH_2O)_n^{-1}.$$

Phenyl. The group C₆H₅-or Ph.

Phenylalanine, α-Amino-β-phenylypropionic acid. C₉H₁₁O₂N.

Colourless crystalline, M.P. 283°C and the solid is soluble is water (3%). An essential amino-acid required by animals.

- m-Phenylenediamine, 1, 3-Diaminobenzene. C₆H₈N₂. Colourless crystals, M.P. 63°C. Basic, diazotization forms brown azo-dyes (Bismark brown). Prepared by reducing m-dinitrobenzene with iron/HCl.
- o-Phenylenediamine 1, 2-Diaminobenzene, C₈H₈N₂. Brown yellow crystals, M.P. 103-104°C. Used as a photographic developer as its solution reduces Ag⁺ ions to Ag. Used for characterising α-diketone,
- P-Phenylenediamine, 1, 4-Diaminobenzene. C₆H₈N₂. White crystalline M.P. 267°C. Cannot be diazotized with HNO₂. Oxidized to quinone derivatives. Its derivatives are used in photography as developer.
- Phenylethanoic Acid. PhCH₂COOH. M.P. 76°C, B.P. 266°C. Occurs in nature as esters. Obtained by hydrolysing benzylcyanide. Used as a penicillin precursor (aqueous solution).
- 2. Phenylethyl Alcohol, PhCH₂CH₂OH. B.P. 220°C, occurs free or combined in many essential oils. (e.g. orange juice). Prepared by Friedel-Crafts reaction between C₆H₆ and ethylene oxide. Used in perfumery.
- N-phenylglycine, C₈H₉NO₂. White crystals, M.P. 127°C. Prepared by heating aniline, NaOH, methanal and KCN. Used to form indigo via indoxyl.
- Phenylhydrazine, C₆H₈N₂, PhNHNH₂. A colourless retractive oil which readily turns brown an exposure to light. B.P. 240°C—241°C. Manufactured by reducing benzene diazonium chloride with SnCl₂ or by treating C₆H₅N₂Cl with Na₂SO₃ (100°C) with subsequent action with HCl and then NaOH.

Subsequent action with
$$1.50^{\circ}$$
 HCl
$$C_6H_5N_2+Cl^-+4H \xrightarrow{Na_8SO_3} C_5H_5NH-NH_8 \xrightarrow{HCl} C_6H_5NH.NH_8HCl$$

Dissolves in organic solvents. Phenylhydrazene is basic fin nature and forms well defined salts, e.g. C₆H₅NH₃+Cl⁻. It reduces Fehling's solution to form a red precipitate. Forms phenylhydrazones and osazones which are characteristics of carbonyl compounds (>CO). Forms antipyrene

Used as a mild oxidizing agent, as a reagent for the identification of aldehydes and Ketones.

Phenylhydrazones. Compounds containing the group Ph NH—N=C=Prepared from carbonyl compounds and phenylhdyrazine. Solids having characteristic melting points. Used to prepare derivatives of indole and for the identification of carbonyl compounds.

with pungent smell, M.P. -33°C, B.P. 162°C. Used as a dehyplus phosgene in HCl.

Phenytion, 5, 5,-Diphenylhydantion, $C_{15}H_{12}N_2O_2$. Used as its sodium phenytoin. Used medicinally in the treatment of focal

Pheromones. Substances secreted by individuals (externally) and which get a specific response from other individuals of the same species, e.g. 'Queen substance' of the honey bee.

Philosopher's Stone. A substance by means of which base metels. like iron could be converted into gold.

Phloroglucinol, 1, 3, 5, -trihydroxybenzene, C₆H₆O₈.

Dihydrate colourless crystals, MP. 200—219°C loses water of crystallization at 100°C. Soluble in water, alcohol and ethersolution. Occurs in many natural glycosides (fuse resorcinol and adhesives.

PH Meter. An instruments used to measure pH of solutions, suspensions, etc. Using glass electrodes. "Null type" and "directreading" instruments are available.

Phorone, di-Isopropylidene acetone, 2, 6,-dimethyl-2, 5-heptadiene-4-one, C9H14O. A yellow liquid with camphor like odour; M.P. 28°C, B.P. 198.5°C. Used as a solvent for cellulose nitrate.

Dimethylmethyl-2-Crotonylphosphate, C7H13O6P. An Phosdrin.

Phosgene, Cl₂CO. A gas possessing unpleasant odour, liquefied at 8°C and is decomposed by water (COCl₂+H₂O →2HCl+CO₂). Prepared by passing Cl₂ and CO(1:1) over charcoal. Also obtain tained by CCl₄ and forming sulphuric acid (CCl₄+2SO₃→COCl₂ +S₂Cl₂O₅). It is also a by-product when chloroform is exposed

Phosphagen. Some creative phosphate present in vertebrate muscles that breaks down on contraction and is resynthesized during the

recovery process.

Phosphatases. A group of enzymes that hydrolyse the several different combinations of phosphoric acid with organic substan-

Phosphate Coatings. Coatings applied to steels and other alloys to make them anticorrosion and render the surfaces good for pain-

Phasphate Esters, (RO)₃PO. Obtained from phosphoric acid, chiefy, H₃PO₄ and alkyl or aryl alcohols, e.g. [(BuO)₃PO, RNA Phosphates. Salts of phosphorus oxyacids, generally, of H₃PO₄,

Phosphatides, Phospholipin (e)s, Phospholipid(e)s. A group of substances of a fatty nature which are essential components of animal and vegetable cells. They contain P and N. Phosphides. Compounds of P with more electropositive element,

e.g. Na₃ P (hydrolysed to PH₃ by water).

Alkyl derivatives of HOP (O) H₂.

Phosphine, PH₂. A colourless gas, slightly soluble in water and has a peculiar fishy smell (water plus calcium phosphide or yellow phosphorus plus conc. alkali). It ignites spontaneously in air. phosphorus plus conc. alkali) oxides of phosphorus. Precipitates Burns in O₂ or air to form oxides of metal salts. Forms phosphorus. phosphides from solutions of metal salts. Forms phosphonium

salts. Hydrogen atoms of phosphine can be replaced by alkyl group (analogous to ammonia).

Phosphinites. Derivatives of R₂P (H) O.

Phosphites. Derivatives of phosphorous acid represented by formula (RO)₈ P. Prepared from PCl₃ and ROH or Ar OH. Used as intermediates in organic synthesis.

Phospholipases. Enzymes that carry out the cleavage of phosphatides either at the carboxylate at the phosphate linkages.

Phospholipids, R'C (O)O CH₂. CH (OC (O) R"). CH₂OP (O)₂OR". A group of substances having a fat-like nature which are essential components of living cell. Lipids are P and N compounds consisting of pounds consisting of number of different species usually with glycerol as 'back bone'. Lipids gets hydrolysed to phosphoric acids and bases. For example, lecithins, cephalins etc.

Phosphomolybdates. A group of salts of heterophosphomolybdic acids mainly of H₃PMo₁₂O₄₀. Used for the estimation of P or

Phosphonates. Compounds with general formula RP (O) (OR)2. Phosphonites. Compounds with general formula RP (OR)2.

Phosphonitriles, Phosphazenes. A group of polymers represented by the formula (PNV) by the formula (PNX₂)₄. Most derivatives have cyclic structures (Polymers). Obtained by heating PCl₅ and NH₄Cl.

Phosphonium Salts. White crystalline salts analogous to the ammonium halides on DILL (DIT) ammonium halides, e.g. PH₄I (PH₃+dry HI). Dissociate into the PH₃ and HX. Triphenylalkyl phosphonium salts with the general formula [(C₆H₅)₃ PCHR/R²]+X⁻ are used for the forwittig reaction.

Phosphoproteins. A type of proteins rich in Pe.g. casein. Forms

Phosphor. A substance that shows luminescence or phosphores-

Phosphoranes. Derivatives of 5-covalent phosphorus, i.e. R₅P,

Phosphoresence. Some molecules after being excited by absorption of visible or ultraviolet radiation and of visible or ultraviolet radiation emit the radiation slowly and long after the light source is removed. This is known as 'phosphorescence' e.g. ZnS, alkaline earth sulphides, etc.

Phosphoric Acids. Oxyacids containing phosphorous (v). Obtained from phosphates and H₂SO₄ or P₂O₅ hydrolysed to H₃PO₄. Phosphates are the salts of phosphoric acid. Used as fertilizers such as superphosphates.

Phosphorimetry. A non-routine technique involving the use of

Phosphorus Acid. Oxyacids of P (III). (Orthophosphorous acid, H₃PO₃ or phosphonic acid). A colourless deliquescent solid (water+P₂O₃ or PCl₃). A dibasic acid producing the anions H₂PO₃ and HPO₃² in water. Forms PH₃ and H₃PO₄.

Phosphorus, P. At. No. 15, At. Wt. 30.973, M.P. (white) 44.1°C, B.P. (white) 280°C, D 1.82. A non-metallic element of group V with electronic configuration 3s² 3p³. Occurs in nature as apatite and fluoroapatite. Prepared from Ca₃ (PO₄)₂ by fusion with and and coke in an electric are furnace, P distils out and is condensed with water. The important allotropes of P are:

(1) White (or yellow) phosphorus (stored under water) and con-

(2) Red phosphorus which is obtained by heating white phosphorous for long at high temperature insoluble in organic A polymeric material.

(3) Black phosphorus which is obtained by heating white phosphorous under high pressure (graphite like structure) with

Phosphorous has high I.P. The stable oxidation states are +5, e.g. in PF₅ and +3, e.g. in PH₃. Forms P_4O_{10} (commonly known as phosphorus pentaoxide, P_2O_5) when burnt in air (excess and P_2O_5) in limited supply of air). Both these (excess and P₂O₃ (in limited supply of air). Both these oxides form acids with water. Also forms ionic phosphides with several metals and metalloids, e.g. Na₃ P or Ca₃ P₂.

Phosphorus compounds are used as fertilizers, in matches, as pesticides, in alloys, special glasses and chinaware, in electrical

Phosphorus (III) Bromide, PBr₃. A colourless liquid (P plus Br₂). Used in organic chemistry for replacing -OH by Br.

Phosphorus (V) Bromide, PBr₅. A yellow crystalline solid which sublimes (PBr₃+Br₂). Hydrolysed to phosphoric acid. Used

Phosphorus (III) chloride, PCl₃ (Phosphorus trichloride). A colourless liquid (P+Cl₂). Hydrolysed to phosphoric acid.

Used in organic reactions to replace -OH by Cl. Phosphorus (V) Chloride, PCl₅ (Phosphorus pentachloride). A white solid (sublimes) formed by PCl₃ plus Cl₂. Used as a chloris

osphorus (III) Chloride Oxide, A colourless liquid (PCl₃ plus chloride, phosphoryl chloride). Phosphorus

O2 or by distilling PCl3 with KClO3) which forms H3 PO4 an hydrolysis. The chlorine atom can be replaced by alkyl groups using Grignards reagent.

Phosphorus Hydrides, PH3 (Phosphine).

Phosphorus, Organophosphorus Acids. Organophosphorus compounds are considered to be the derivatives of phosphorus

Phosphorus (III) Oxide, P₂O₃ (Phosphorus trioxide). A white waxy solid with a garlic smell, usually exists as P₄O₆ molecule. Soluble in organic solvents (e.g. CS₂, C₆H₆), Obtained by burning P in a limited supply of air. Oxidises to P2O5 by air at room temperature and inflames above 70°C. Forms phosphoric acid or one of its salts with cold water or dilute alkalis and POCl3 with Cla.

Phosphorus (V) Oxide, P2O5 (Phosphorus pentoxide). A white powder soluble in organic solvents. Exists as P₄O₁₀ molecules. Obtained by burning P in excess of air. Forms phosphoric (V) acid on the hydrolysis. Used as a dehydrating agent because it removes water from some oxy-acids and other oxygen contain-

Phosphorus Oxyacids. Hypophosphorous acid, H₃PO₂, H₂P(O)OH. A monobasic acid. The acid and its salts are strong reducing

Phosphorus Acid, H₃PO₃, HP (O) (OH)₂. A dibasic acid-a strong reducing agent. Obtained from PCl₃ and H₂O Hypophosphoric Acid, H₄P₂O₆. HP (O) (OH) (μ -O) P (O) (OH)₂. Orthophosphoric Acids, H₃PO₄, OP (OH)₃. A tribasic.

Pyrophosphoric Acids, H₄P₂O₇, (OH)₂ P(O) (μ -O) P(O) (OH)₂. Phosphorus Polysulphides. Yellow solids made up of P and S

e.g., P₄S₃, P₄S₄, P₄S₅, P₄S₇ and P₄S₁₀. Obtained from red phosphorus and sulphur. Used in the manufacture of matches.

Phosphorylase. An enzyme that facilitates the addition of phosphate across glycosyl or related bonds. Widely distributed in plant and animal tissues.

Phosphotungstic Acids. A group of fungstion complex acids, e.g. H₇[P (W₂O₇)₆]. The anions of the acid contain P, W and oxy-

Photoacoustic Spectroscopy. A spectroscopic technique involving the irradition of the sample by chopped monochromatic radiation and the absorption of the radiation causes a temperature fluctuation which is transmitted to the gas in contact with the sample and which in turn transmitted to microphone. Used to measure the optical absorption spectra of solids and liquids

Photochemistry. The investigation of chemical changes which occur in consequence of absorption of photons from the visible

and ultraviolet radiations having wnvelengths approximately between 10,0000A and 1000A and the reactions are called photochemical reactions. The absorption of light is a quantum process. The energy in this range varies from 3,00,00 cal to 28,600 cal per mole. For example, decomposition of HI, photochemical combination of H2 and Cl2, etc.

- Photoconduction. A process in which some substances such as selenium, which are poor conductors of electricity in the dark, become good conductors when exposed to light (transfer of electron from valence band of the solid into the conduction band).
- Photodissociation. A photochemical reaction in which a molecule dissociates into smaller molecules, radicals or atoms, e.g. dissociation of a molecule of acetone by absorbing a quantum of light.

(CH₃)₂ CO→2CH₃+CO· methylradical.

Photo-Electric Cells, Photoelectric Effect. The emission of electrons from a solid (or liquid) surface, mainly the alkali metals, when it is irradiated with electromagnetic radiation (of suitable wave length). This phenomena is known as photoelectric effect. the photoelectric effect, the number of electrons emitted depends on the intensity of the radiation and not on its frequency and this fact can be made the basis of an instrument for measuring light intensity (photoelectric cell). To remove an electron from the solid a certain minimum energy must be supplied, known as work function ϕ so there is a minimum threshold ϕ threshold frequency vo for radiation to eject electrons: $hv_0 = \theta$. If the frequency is higher than this threshold energy, the elec-

Photo-Electron Spectroscopy. When an atom is bombarded with photons of suitable energy, the electron of energy (E) is emitted and $E=E_i-E_b$, where E_i and E_b are the respective energies of photon and of binding of the ejected electrons. This technique is used for measuring binding energies of valence electrons or for inner core electrons. This is known as inner core spectroscopy or electron spectroscopy for chemical analysis (ESCA) or X-ray photo-electron spectroscopy (XPS).

Photoelectron. An electron ejected from a solid, liquid, or gas by the photoelectric effect or by photoionization.

Photoemission. The emission of photoelectrons by the photoelectric

Photographic Developers. Chemicals used to reduce silver halide grains in the photographic plate which have been exposed to

Photographic Gelatin. A protective colloid medium for silver halide emulsion used in photography.

Photoluminescence. A general term applied to the emission of light as a result of an initial absorption. For example, fluorescence

Photolysis. A-chemical reaction produced by electromagnetic radiation (light or UV radiation). Many photolytic reactions involve the formation of free radicals.

Photon. A quantum of light energy. The energy of a photon is given by $E = hv_0$, where h is Planck's constant and v is the frequency of radiation

Photosensitization. A process in which a suitable foreign matter when mixed with reactants absorbs the radiation and becomes excited and subsequently, pass on this energy (absorbed) to the reactant molecules which will undergo dissociation or chemical transformation. Such reactions are termed photosensitized reactions and the foreign substance which absorbs and transfers the energy of radiation is called photosensitizer. Mercury and cadmium vapours are often used as sensitizers in reactions such as dissociation of hydrogen by a light of wavelength 2536 A.

Photostationary State. A state in a phochemical reaction when the rate of removal of reactants by light is equal to the rate of recombination of reaction of the products to yield the reactants.

Photosynthesis. A process occurring in plants and by which they build up their carbon compounds from atmospheric carbon dioxide using light as the source of energy.

Phototropy. A term denoting the reversible change of colour which certain organic substances undergo when exposed to light of a suitable wave length. The original colour returns in the dark, and the reversion is accelerated by heating.

Phthalamide, C₈H₈N₂O₂, C₆H₅ (CONH₂)₂. Colourless crystals, M.P. 200.210°C (decomposition) very slightly soluble in water. Obtained by the action of cold concentrated and acquired acquired to the action of cold concentrated and acquired acquired to the action of cold concentrated and acquired acquired to the action of cold concentrated and action of cold concentrated and acquired to the action of cold concentrated and acquired to the action of cold concentrated and action of cold concentrat Obtained by the action of cold concentrated ammonia solution

Phthalic Acid, C₈H₆O₄,

Colourless crystals,

M.P. 190-210°C (decomposes). A dibasic acid oxidation of naphthalene or o-xylene) which forms stable salts and benzene

Phthalic Anhydride, C₈H₄O₃ Long needles. M.P.

130°C, B.P. 284°C. Prepared by oxidizing naphthalene.

$$\frac{270 - 300^{\circ}\text{C}}{\text{H}_{2}\text{SO}_{4}/\text{HgSO}_{4}}$$
CO
$$\frac{\text{CO}}{\text{CO}}$$

or by vapour phase oxidation of naphthalene over V₅O₅ (commercial method). Used to prepare substituted anthraquinones, phthalein dyestuffs, quinoline dyes, etc.

Phthalide, C₈H₆O₂ M.P. 75°C. Obtained by reducing phthalic

anhydride with Zn dust and NaOH.

Phthalimide, C6H5NO2,

less plates (ammonia passed in molten phthalic anhydride). Reduced to phthalide with Zn/NaOH solution. Hydrolysed to phthalic acid or its salts by prolonged hydrolysis with acids or alkalis. Used in Gabriel synthesis.

Phthalocyanines. A type of organic colouring matters generally used as pigments and more limited as dyestuffs. They show great fastness and brilliancy of shade (green to blue). Obtained by, say, heating phthalic anhydride with urea (or ammonia) and a metallic salt. Phthalocyanine molecules is reasonably stable to heat and chemical reagents and is formed by the union of of one atom of the metal with four molecules of phthalonitrile, e.g. copper phthalocyanine (C₃₂ H₁₆ Cu N₈).

Phthioic Acids, Phthienoic Acids. A group of branched-chain fatty acids isolated from mycobacteria (C22 to C28).

Phytanic Acid, 3,7,11,15,-tetramethyl hexadecanoic Acid, C₂₀H₄₀O₂.

Phytic Acid, C₆H₆(O(PO) (OH)₂)₆ (Meso inositol hexaphosphoric acid). A syrupy liquid, soluble in water. Occurs in seeds as insoluble calcium magnesium salt. Also found in blood plasma.

Phytochemistry. The chemistry of compounds which are present in

Phytol, 3,7,11,15 tetramethyl-2 hexadecen-1-01, C20H40O. Colourless, oily, diterpenic alcohal, B.P. 202-204°C.

 $P_i(\pi)$ bonding. See orbitals.

- Pickling. The process of immersing materials in a bath of either dil H₂SO₄ or HCl for a short period, followed by washing in water. The process is carried out to get a clean scale-free surface, e.g. a surface of iron sheet before galvanizing.
- Picolines, C₆H₇N. Methyl pyridines found in bone oil and coalfar $(\alpha, \beta \text{ and } \gamma \text{ picoline})$.

Picramic Acid,
$$C_6H_5O_5N_8$$
 O_2N O_2N O_2N O_2N O_2N O_2N O_2N O_2N O_3N O_4N O_2N O_2N O_3N O_2N O_3N O_3N

Soluble in hot water, obtained from picric acid reduction with sodium hydrogen sulphide. Used to prepare 930-dyes.

Picric Acid, 2-amino-4,6-dinitrophenol. O2N NO2

Lemon yellow prisms, M.P. 122.5°C, soluble in alcohol-Obtained by nitrating phenol. In the cast form also known as lyddite. Forms characteristic derivatives with many hydrocarbons. Used for dyeing wool and silk.

- Picrolonic Acid, C₁₀H₈N₄O₅, M.P. 124°C. Fine yellow needles. Used for the isolation and identification of organic bases, and in tests for calcium, copper and lead.
- Piezoelectricity. It is the electric charge developed in anisotropic crystals when they are subjected to stress. Used in electrical
- Pig Iron. The crude iron produced from a blast furnace, containing C, Si and other impurities. The molten iron is allowed to run out of the furnace into channels (called 'sows'), which branch out into a number of offshoots (called 'pigs') in which the metal is allowed to cool.
- Pigments. An insoluble coloured material used to impart colour to surfaces, plastics, in inks, etc. Many pigments are inorganic, such as. TiO₂, ZnO, ZnS, lead, cadmium chromates, ultramarines, etc. Some dyestuffs are also used as pigments.
- Pimelic Acid, Heptanedioic Acid, C₇H₁₂O₄, HOOC. [CH₂]₅ COOH M.P. 105°C. Colourless crystals obtained by oxidizing capric acid or oleic acid. Used in polymer chemistry.
- Pinacol, Pinacone, 2-3-dimethyl-2, 3-butancdio C₂H₁₄O₂. M.P. 38°C B.P. 175°C colourless crystals. Obtained by reacting benzene and propanone mixture with magnesium amalgam. Crystallizes out as hexahydrate (M.P. 47°C) water. Forms pinacolone.
- Pinacolone, Pinacolin, 3, 3 dimethyl-2-butanone C2H12O. Colourless liquid, B.P. 103-106°C at 746 mm. Smell like camphor.

Pinacol-Pinacolone Rearrangement. When heated with minerals or organic acids, pinacol undergoes a molecular rearrangement with loss of water to form pinacolones (ketones).

ss of water to form pinacolones (ketones):

$$CH_3 CH_3$$
 CH_3
 $CH_$

Pinacols. 1, 2-glycols of the type

where R₁, R₂, etc., are alkyl or any radicals.

Pinane. C10H18

A saturated hydrocarbon obtained by the catalytic hydrogenation of α or β pinine. Exists in cis and trans forms.

a-Pinene, C₁₀H₁₆. A dicyclic terpene, B.P. 156°C. Present in most essential oils, derived from coniferae and a main constituent of turpentine oil. Obtained from turpentine oil (+ form). Forms

pinane when reduced in the presence of a catalyst; used to form artificial camphor (HCl gas passed through turpentile oil).

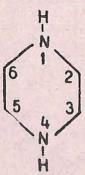
β-Pinene (Nopinene). C₁₀H₁₆, (+)-β-Pinene occurs only one in oil, extracted from the ripe fruits of Ferula galbaniflua. B.P. 162-

Pink Salt. Ammonium hexachlorostannate.

Pino camphone, C₁₀H₁₆O. A dicyclic ketone. Found in oil of hyssop in the (-)-form (B.P. 211°C).

Pinocarveol, C₁₀H₁₆O. A dicyclic terpene found in the oil of Euclayptus globulus((-)-form).

Piperazine, Hexahydropyrazine, C₄H₁₀N₂. Colourless crystals,



M.P. 44°C (hydrous) 104°C (anhydrous) obtained from alc ammonia and 1, 2, dichloroethane. Used to treat threadworms

A colourless liquid_r. Piperidine, Hexahydropyridine, C,H11N.

miscible with water, M.P. -, °C, B.P. 106°C with a smell of ammonia. Obtained by heating peeper with alkali. Also prepared by reducing pyridine. A strong base analogous to aliphatic amines.

Piperine, C₁₇H₁₉NO₃. An alkaloid obtained from pepper.

Piperitol, C₁₀H₁₈O. An optically active, s-terpene-alcohol. Occursin (-) form in eucalyptus trees. A pleasant smelling viscous liquid.

CH₂

Piperitone, C10H16O. An optically active terpene ketone, occurs in eucalyptus oil and peppermint oil. Colourless oil, B.P. 109.5-110°C.

Piperonal, C₈H₆O₃. White crystals M.P. 37°C, B.P. 263°C. Occurs in association with vaniline. Prepared by oxidizing piperine, Used to perfume soaps.

Pipette. An apparatus used to measure volume of solutions (in volumetric estimations).

Pitchblende. UO_{2-2.5}. An ore of uranium.

Pitch, Coal Tar Pitch, A residue of coal-tar distillation.

Pivalic Acid, 2, 2-Dimethyl Propanoic Acid, (CH₃)₃.CCOOH. Colourless solid, M.P. 35.5°C. Prepared by the carboxylation of isobutene with CO/H2SO4.

Pk. The negative logarithm of the equilibrium constant of an equilibrium reaction.

Planar Complexes. Describing complexes in which the acceptor atom is surrounded by four ligands in a plane, e.g., [Pt(NH₈)₄]²⁺

Planck's Constant, h. A constant used in the equation, $E=hv_{s}$. where v is the frequency of radiation. It has a value of 6 6256 × 10-27 erg sec.

Plane of Symmetry. See Symmetry elements.

Planetary Electrons. Extranuclear electrons of an atom.

Plant Hormones. Compounds responsible for physiological processes in plants, e.g., auxins.

Plaster. A material containing (I) a mixture of lime and sand or of portland cement and sand, and (2) materials composed of plaster of paris.

Plasma. A mixture of ions and electrons as in an electric discharge.

Plaster of Paris. A partially hydrated calcium sulphate, 2CaSO₄, H₂O, formed by heating gypsum (CaSO₄, 2H₂O). The setting of the plaster takes place due to rehydration by water.

Plastic Explosives. Describing a type of explosives which can be moulded by hard, e.g., cyclonite mixed with an oil.

Plasticizers. High molecular weight liquids or low melting solids incorporated into plastic resins to change workability, flexibility, flow and impact resistance. E.g. halogenated hydrocarbons,

Plastics. Polymeric materials, organic in nature and shaped by flow. They are artificial and are classified as:

Thermoplastics—Softned by heating and rehardened on cooling e.g., polyethene.

Thermoset or Thermocured plastic. Infusible and insoluble in their final moulded shape final moulded shape, e.g. phenol fermaldehyde resins.

Plastoquinones. A group of trisubstituted benzoquinones having polyisoprenyl groups. Occur in chloroplasts and used in photo

Plate, Tray. A tray used in fractionating colums.

Plate Column, Tray Column. A distillating column consisting of a large number of perfect. A distillating column large number of perforated horizontal plates (equally spaced).

Plate Efficiency, Tray Efficiency. The distance between successive plates in a plate or bubble. plates in a plate or bubble cap distillation column. If a plate in a distillation or absorption a distillation or absorption column were 100% efficient, the liquid and vapour streams 1 column were 100% efficient, aquililiquid and vapour streams leaving the plate would be in equili-

Platinum, Pt. At. No. 78, At. wt. 195'09, M.P. 1772°C, B.P. 3827°C D2!.45. A transition metal and property of the control of t D2!.45. A transition metal occurring naturally and in traces in heavy metal sulphide ores. heavy metal sulphide ores. It is the heaviest element of the nickel group, electronic conf. nickel group, electronic configuration 5d⁹6s¹. Shows oxidation states of +6 and +5 [fluoristation 5d⁹6s¹. Shows oxidation states of +6 and +5 [fluoride derivative only), +4, +2, +1 and 0. It is very malleable and +1 and +1 is very malleable and +1 is +1 in +1 and +1 is +1 in +1and 0. It is very malleable and ductile, resistant to oxidation and is not attacked by acids (except. aqua regia) or alkalis. Attacked by F. and Cl. above 2000 aqua regia) or alkalis. Attacked by F₂ and Cl₂ above 300°C. Pt(II) and Pt(IV) form a series of complexes (square 1) series of complexes (square planar or octahedral co-ordination

respectively). The metal is used in jewellery, laboratory ware, thermocouples, electrical contacts and as a catalyst (for synthesisof NH.).

Platinum Ammines. Co-ordinate complexes containing Pt-NH3

(and amine) groups.

Platinum Black. A finely divided precipitate obtained from Pt(II) solutions by reducing agents. Used as a hydrogenating catalyst.

Platinum Halides:

Platinum fluorides. PtF2 and PtF4 are formed when metal is heated in F2.PtF4 is hygroscopic (soluble in water) while Pt F2 is yellowish green and insoluble in water.

Platinum iodide. PtI2 (black powder) and PtI4 (black powder)

are obtained by double decomposition.

Platinum chloride. PtCl4 (red-brown), PtCl3 (green-black) and PtCl₂ (red-black) are all formed from Pt and Cl₂.

Platinum hydrosol. A powerful reducing agent prepared by reducing platinum solutions with, e.g., hydrazine hydrate in presence of a protective colloid.

Platinum Iodides Complex. Iodoplatinic acid, H2PtI6, 9H2O. Prepared (red monoclinic crystals) from PtI4 in hydriodic acid. A number of salts such as (NH₄)₂ PtI₆ and K₂PtI₆ and known.

Platinum Iridium. An alloy of Pt containing up to 35% iridium. Its hardness and resistance to chemical attack increases with the increase in iridium content. Used in jewellery, electrical contacts and hypodermic needles.

Platinum-Metals. The group of transition metals, ruthenium (Ru), osmium (Os), rhodium (Rh), iridium (Ir), palladium(Pd), and platinum (Pt), They occur together as native alloys, in mixed sulphide ores and as traces in Au, Ag and Cu sulphide ores.

Platinum Organic Derivatives. Stable alkyl and aryls, e.g, [trans-PtBrMe(PEt3)2] formed by the action of Grignard's reagents on

Platinum Oxides PtO₂ (brown) formed by dehydration of the hydrated oxide precipitated on hydrolysis of Pt Cl₆²⁻. PTO (PtCl₂+KNO₃) and Pt₃O₄ are also known.

Pleochroism. Discribing the property of a crystal of having a differrent colour depending upon the direction of transmitted light through the crystal, e.g. K2Pt(CN)4.

Plumbates. (Lead (IV) hydride, PbH4). A colourless unstable gas obtained by the action of acids on a mixture of Mg and lead

pellets.

Plumbates. See lead oxide.

Plumbic. Lead (IV) Compounds.

Plumbous. Lead (II) Compounds.

Plutonium, Pu. At. No. 94, M.P. 641°C, B.P. 3232°C D 19'82. A highly toxic radioactive silvery element of the actinoid series of metals. A transuranic element found on earth only in minute quantities in uranium ores but readily obtained, Pu-232, by neutron bombardment of natural uranium. Pu is separated by selective oxidation and solvent extraction. The metal is prepared by reduction of PuF₄ with Ca. Used as a nuclear power source (i.e., in space exploration).

Plutonium compounds. The metal reacts similar to that of U, slowly with water and rapidly with dilute acids. In aqueous solutions the +3, +3, +4, +5, and +6 oxidation states co-exist. PuF₆

red-brown (PuO₂+HF+O₂). PuO₂ is a stable oxide.

Plywood. Layers of wood bonded by adhesives, e.g. a laminate.

Pnictogens. A name given to the elements nitrogen, phosphorous, arsenic, antimony and bismuth of Group V. They are characterised by the presence of X3- species.

Po. Polonium

Point Group. A set of symmetry elements passing through or referring to single point.

Poison. A substance that destroyed catalyst activity.

Poiseuille, Law of. A law describing the flow of a liquid through a capillary.

$$\eta = \frac{\pi P r^4 t}{8l V}$$

where η =the coefficient of viscosity of the liquid, P=the pressure head of the liquid, r=the radius of the Capillary l=length of the tube, V=volume of the liquid flown through the tube in time T.

Polar. Describing a compound with molecules that have a permanent, dipole moment e.g. HCl and H₂O.

Polar Bond. See covalent bond.

Polarimeter. An instrument used to measure the rotation of plane polarized light.

Polarizability. When a molecule is placed in an electric field the electrons tend to be drawn away from the nuclei with the result that a dipole is induced in the molecule.

Polarizability of the molecule The strength of the dipole

Electronic Polarizability and atomic polarizability are also known. Polarizability has units of volume.

Polarization. The restriction of the vibration in a transverse wave so that the vibration occurs in a single plane. The important polarizations are as follows:

Orientation polarization. In this polarization dipole molecules when placed in an electric field try to align themselves in a certain direction within the field.

Electron polarization. Electrons within each molecule are displaced slightly in the direction towards the +ve pole of the field. Atomic polarization. The nuclei having a +ve change are slightly displaced relative to each other.

Molecular polarization=Orientation polarization+electronic polarization+atomic polarization.

- Polar Molecules. A molecule constituted by atom of different electronegativities and by a covalent link, e.g. H8+-C18molecule.
- Polarography. An analytical method in which current is measured as a function of potential and particularly used for dilute solutions of substances which are susceptible to electrolytic reduction at a mercury cathode. A dropping mercury electrode (DME) is used to record polaragrams (current voltage curves). The method is used for determining small amounts (traces) of metals, complexes in metals and for non-aqueous solutions.

Polar Solvents. Solvents with good solvating powers, greater dipole moments which act as better solvents for ionic species.

Pollucite, Cs (Al Si₂ O₆), xH₂O. An impotant mineral of Cs.

Polonium, Po. At. No. 84, At Wt. 210, M.P. 254°C, B.P. 962°C. A radioactive decay product from heavier elements belonging to group VI of periodic table. Electronic configuration 5d10 6s2 6p4. Prepared by nentron irradiation of 209Bi. It is an α-emitter with over 30 radio isotopes. Po salts are generally coloured (yellow or red).

Polonium Halides. PoX4 and PoX2 are well known halides. PoCl2 and PoBr2 are formed by the thermal degradation of the tetra-

Polonium Oxide, PoO2. A basic oxide obtained from Po and O2 at 250°C.

Poly. Polymeric derivative.

Polyacetals. Polymers obtained from polyols, e.g. HO (CH2)n OH and carbonyl derivatives, e.g. CH₂O. Polyacetals such as HO [(CH₂)ⁿ OCH₂O]_x (CH₂)ⁿ OH one obtained from polyols and carbonyl derivatives. Used in coatings.

Polyacrylamide, CH2: CHCONH2. Linear polymers with high molecular weight, gels, gums, plastics soluble in water.

Polyacrylonitrile. Important fibre elastomers (CH2 CH CN)n. Used as a co-polymer with, e.g. butadiene, styrene, etc.

Polyalkanes. Polymeric olefins.

Polyalkylidenes, (CHR) A group of polymers. Prepared from diazomethane and alcohols. Their properties depend upon the degree of polymerization and the stereochemistry,

Polyamides. A synthetic polymer in which the monomers are linked by-NH-CO-group. E.g. nylon.

Polyamines, -(HN-R-NH-R'), Hydrophilic polar substances used as flucculants for cellulose fibre, etc.

Polybenzimidazoles. Polymers containing benzimidazole rings. Prepared by condensing aromatic diamine and dicarboxylic acid derivatives in the presence of P2O5 in H3 PO4 or high boiling point solvents. Used in adhesives and laminates.

Poly Blend. A mixture of polymershaving of different structures.

Polycarbonates. A class of thermoplastics represented by the general formula

H (OROC), OROH

and show toughness, high softening point and clarity.

Polychloral. An aldehyde polymer.

Polychloroethylene. See tetrachloroethene (Polyvinyt chloride PVC). A synthetic polymer obtained from chloroethene.

Polychloroprene. See neoprene.

Polycrystalline. Describing a substance constituted of a large number of many interlocking crystals that have solidified

Polycyclic. Compound having two or more rings in its molecule.

Polyene. An alkene with more than two double bonds in its mole-

Polydentate Ligand. See multidentate ligand.

Polydispersion. Sols in which the dispersed particle sizes vary

Polyelectrolyte. A macromolecular compound containing many ionizable groups within the same molecule. There may be purely anionic, cationic or amphoteric polyelectrolytes, e.g. polystyrene sulphonic acid (weakly acidic), polymethacrylic acid (strongly basic), polyvinylamine (amphoteric), etc.

Polyene Antibiotics. Macrolides.

Polyenes. Describing compounds having many carbon to carbon

Polyester. A synthetic polymer formed by condensation of polyhydric alcohols and polybasic acids, e.g. glycol or propylene glycol and maleic or terephthalic acid.

$$-\left(CH_{2} CH_{2} OC(O) - \left(\begin{array}{c} \\ \end{array}\right) - C(O) - O \right)_{n} - C(O) - O = C(O) -$$

For example, synthetic fibres such as Terylene and Dacron are

polyesters.

Polyethene (Polyethylene, polythene). A synthetic polymer made from ethene (Ziegler's method). Formed in soft and hard varieties.

Polyethers. Polymers containing (C-O-C)n units in the back-

bone, e.g. aldehyde polymers.

Polyethylene Glycols. HO (CH2 CH2 O)n H. High molecular weight polyether glycols obtained from ethylene oxide and water, dihydroxyethene or diethylene glycol plus base. Used as solvents and in pharmaceuticals.

Polyhalides. Describing compounds containing complex amions (halogens and interhalogens plus halides), e.g. [ICIBr].

Polyhexafluoropropene. Fluorine containing polymers.

Polyimides. Compounds containing the imide group-C (O)-NC (O) -, e.g. film, plastics and wire enamels. They have good heat-resistant properties.

Polyiodides. Specific polyhalides containing species such as

Is, I,, I,.

- Polyketides. Naturally occurring organic compounds considered as arising from poly-β-ketonic intermediates derived by condensation of ethanoate units. Biosynthesis of some reactions such as fatty acids, tetracyclic antibiotics, etc. takes place via polyketide routes.
- Polymeric Reagents. Reagents which are solids, insoluble and used for organic synthesis. Their action is selective, simplified and high purity product isolation. Obtained by introducing organic functional group such as Br, Li, SO₃ H etc. on to a small proportion of benzene rings of cross-linked polystyrene.

Polymerization. A process in which one or more compounds react

to form a polymer.

Monomers are formed by the combination of two or more units

of same compound.

Macromolecules are obtained by the repeated structural units. These structural units are known as mers. Addition polymerization occurs when the monomers undergo addition reactions, with no other substance formed, e.g. the formation of polystyrene from styrene:

Ph CH=CH₂—→[PhCHCH₃-]_n. polystyrene

Condensation polymerization. A process in which polymer is formed by reaction of monomers with the elimination of small

molecules, e.g. formation of Terylene by the condensation polymerization of ethylene glycol with terephthalic acid.

$$n$$
HO . $[CH_2]_2$.OH+ 4 HO.CO \bigcirc CO₂H
$$-\left(\begin{array}{c} O.[CH_2]_2.O.CO\bigcirc \bigcirc \end{array}\right)$$
. CO $\left(\begin{array}{c} - + nH_2O \end{array}\right)$

Polymethine Dyes. Dyestuffs having two polar atoms joined by a methine chain [Y (CH) $_n$ =X] containing an odd number of atoms in the chain.

Polymethylene Glycols. Compounds formed by the union of several molecules of formaldehyde with a molecule of water. They are of the type

HO-CH₂OCH₂O.....CH₂OH,

and are formed in aqueous solutions of methanal. The size of the molecule increases with the increase in concentration of

Polymorphism. If a substance is able to exist in more than one crystalline from, it is said to be polymorphous and the phenomena is known as polymorphism. Polymorphism when applied to elements is known as allotropy.

Polymyxins. A class of closely related antibiotics produced by

Poly (oxyphenylenes), Poly (phenylene oxides), Poly (phenylene ethers). Polymers formed by oxidative polymerization of phenols using O₂ with Cu and an amine (pyridine) as catalysts.

$$\begin{bmatrix} -R \\ -R \end{bmatrix} - \begin{pmatrix} R \text{ is generally Me} \end{pmatrix}$$

Thermoplastics so formed are used in electrical appliances.

Polypeptides. See Peptides.

Poly (phenylene sulphides). -(-___S_\mathbb{\text{\subset}}_n

Commercially prepared by the polymerization of S, Na₂ CO₃ and C₆H₅X in a sealed container at 275-370°C. Used as adhesives (high temperature) for laminates and in coatings.

Polyprenols. Primary monohydroxylic alcohols having a carbon skelton and is made up of isoprene units linked head to tail. They are bisynthesised from mevalonic acid. carriers in the transfer of sugars from nucleotide diphosphate sugars to a wide range of acceptors.

- Polysaccharides. High molecular polymers of the monosaccharides or sugars or carbohydrates derived from monosaccharides by the removal of n-1 molecules of water from n molecules of monosaccharides. Imorpant polysaccharides are inulin, starch, glycogen and cellulose. They can be broken down to smaller polysaccharides, disaccharides and monosaccharides.
- Polystyrene. A range of polymers made from styrene which constitutes an important member of the group of ethenoid plastics. One of the lightest ethenoid plastics with negligible water absorption and outstanding electrical properties at high frequencies. Used as rigid insulation in radio applications such as radar and television.
- Polysulphide Polymers. Polymers containing (SRSSRS), groups, obtained from dihalides. They are good elastomers and good solvent resistant. Used in printing rolls and paint spray.
- Polysulphides. Species formed as aqueous solutions from sulphides and sulphur. Contains $[S_2]^{2-}$, $[S_3]^{2-}$, $[S_4]^{2-}$, $[S_5]^{2-}$ and $[S_6]^{2-}$ anions. Used for the manufacture of polysulphide polymers.
- Polyether derivative obtained from bisphenol A and 44'-dichlorodiphenyl sulphone.
- Polytetrafluoroethene, Teflon, Fluon, PTFE. A synthetic polymer made from tetrafluoroethene (i.e. CF2: CF2). A very tough, translucent material with a low coefficient of friction and high softening point (320°C). Used as electrical insulator and as a
- Polythionic Acids, Polythionates, H2SnO6. A class of dibasic acids (more or less unstable) prepared by the action of I2 solution on sodium sulphite, thiosulphate and mixture of these, by the action of H₂S on other members of the same series and of H₂S on H₂SO₃ at 0°C. For example, dithionic acid (H₂S₂O₆), trithionic acid (H₂S₃O₆), tetrathionic acid (H₂S₄O₆), pentathionic acid (H₂S₅O₆) and hexathionic acid (H₂S₆O₆).
- Polyurethane. A synthetic polymer containing the group-NH-CO-O linking the polymer. Prepared from polyhydroxy compounds and polyisocyanates. Used in foams.

Poly (vinyl acetals). A class of resins obtained by the action of poly

(vinyl alcohol) on aldehydes.

cohol) on alderlydes.
$$-(CH-CR_2-CH-CH_2)_n-$$
O
$$-(CH_2)_n$$

Used in metal paints, wood-sealers, adhesives and in safety glass

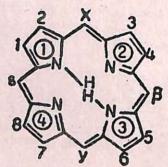
Poly (vinyl alcohol). Polymers containing the unit - (CHOH-CH₂)_n-, Formed by the hydrolysis of the acetate by NaOH and water or

from polyvinyl acetate. Used as a size in the textile industry, in aqueous adhesives, in the manufacture of polyvinyl acetates for safety glasses.

Prophobilinogen, C₁₀H₁₄N₂O₄. An immediates biogenetic precursor of the prophyrins.

Porphyrinogens. A group of naturally occurring pigments containing methylene bridges. They are biosynthetic intermediates which are instantaneously oxidized to porphyrins.

Porphyrins. A class of naturally occurring pigments. Haemoglobin and other animal respiratory pigments and chlorophyll and the respiratory catalyst of plants constitute compounds of porphyrins with metals. They are derived from porphin which is shown below:



Important porphyrins are:

Protoporphyrins, $C_{34}H_{34}N_4O_4$ (present in haem),

Mesoporphyrins, $C_{34}H_{38}N_4O_4$ (from haemin+HI),

Haemoatoporphyrin, $C_{34}H_{38}N_4O_6$ (haemin+strong acid)

Coproporphyrin $C_{36}H_{38}N_4O_6$ (present in urine in the serum of animals and in the feaces)

Portland Cement. A hydraulic cement made by calcinating a mixture of calcium carbonate (CaCO₃) and some aluminium silicates in suitable ratio to produce clinker which is powdered to form usuable cement. The important constituent of portland cement is 3CaO, SiO₃.

Positive Rays or Canal Rays. The ionized gas molecules having positive charge and which are produced in discharge tube when an electric current is passed at low pressure. They move through a cut in the cathode in the form of a beam.

Positive Ray Analysis. When a bean of positive particles is passed through a magnetic and an electric field, the various ions are deflected to different extents, depending on their velocities, masses and charge. This analysis is the basis of a mass spectrometer.

Positron, β+. The positive counterpart of the electron and also an indivisible unit of positive electricity.

Positronium. One electron and one positron (analogous to hydro-

Post-actinide Elements. A group of elements following 103Lw and which should be part of a 6d transition series.

Potash Alum. See alum.

Potassamide (potassium amide), KNH2. White solid obtained from K and NHa.

Potassium, K. At. No. 19, At. wt. 39 09, M.P. 63 65°C, B.P. 774°C, D 0.86. A soft reactive metal, Videly distributed in nature in silicate rocks, e.g. Orthoclase, K Al Si, O8, in plants, in milk, etc. also occurs as carnallite, KMg Cl₃, 6H₂O (in salt beds). The third member of the alkali metals (I group). The electronic configuration is 4s¹. It shows only+1 oxidation state (in simple compounds). Forms complexes with water, NH, (unstable type) and macrocyclic ligands such as crown ethers. Prepared by reducing KCl with Na at high temperature and low pressure or by electrolysis. Forms KO₂ (superoxide) when burnt in air. The important isotope of potassium is ³⁹K (93.1%). Used as a reducing agent, a Na/K alloy may be used for heat transfer. Also used as fertilizers.

Potassium Antimonyl Tartrate (Tartaremetic), C₄H₄O₇ Sb K, ½ H₂O. Colourless rhombic crystals soluble only 6% in cold water, Used medicinally as an emetic.

Potassium Bromate, KBrO₃. Obtained from Br₂ plus hot aqueous KOH. Used as a standard in volumetric work.

Potassium Bromide, KBr. Obtained from KOH or K2 CO3 and HBr or Br₂ plus KOH. M.P. 728°C. Used as a sedative.

Potassium Carbonate Pearl Ash, K₂CO₃. A white deliquescent solid manufactured by Leblanc process. It crystallizes out between 10 and 25°C as K₂CO₃, 3H₂O. It dehydrates at 100°C to K₂CO₃, H₂O and at 130°C to K₂CO₃. Used in salting out alcohol from aqueous solutions, as a standard and in the manufacture

Potassium Chlorate, KClO₃. A white solid obtained by the electrolysis of a concentrated solution of KCl. Manufactured by the fractional crystallization of a solution containing NaClO₃ and KCl. On strong heating forms KCl and O2. When heated just above its melting point, potassium perchlorate (KClO₄) is formed

Potassium Chloride, KCl. Occurs in nature as sylvine and in sylvinite (KCl-NaCl) carnallite (KMg Cl₃, 6H₂O), Kainite (KCl. Mg SO₄, 3H₂O), hard salt (KCl-NaCl-MgSO₄). Prepared by neutralizing HCl with KOH. Used as a fertilizers.

Potassium Chromate, K₂CrO₄. A bright yellow solid prepared from KOH and K₂ Cr₂ O₇ solutions. Soluble in water and addition

- of an acid to its aqueous solution converts the chromate ions (CrO_4^{2-}) to dichromate ions $(Cr_2O_7^{2-})$. Used as an indicator in argentometric titrations.
- Potassium Citrate, K₃C₆H₅O₇, H₂O. Colourless crystals, soluble in water and used in medicines as diuretic.
- Potassium Cyanide, KCN. A white ionic solid, M.P. 635°C and prepared by fusing K₂CO₃ plus C plus NH₃ gas (Beilby's process). An extremely poisonous salt.
- **Potassium Dichromate**, K₂Cr₂O₇. An orange red solid (chromite+ K₂CO₃+CaO followed by acidification) soluble in hot water (more). Used to prepare chrome pigments, as an oxidizing agent in organic chemistry and in volumetric analysis.
- Potassium Ferricyanide, K₃ [Fe(CN)₆]. Dark red crystals obtained by the oxidation of potassium ferrocyanide with Cl₂ and the ferrocyanide is obtained from the mixture by fractional crystal-
 - $2K_4[Fe(CN)_6]+Cl_2\rightarrow 2K_3[Fe(CN)_6]+2KCl$. A powerful oxidizing agent and acts both in acid or alkaline medium. Used
- Potassium Ferrocyanide, K_4 [Fe (CN)₆]. A brown yellow crystalline substance obtained from: (a) crude canstic potash when heated with iron filings with nitrogenous matter (horns). $4KOH+Fe+8C+8N\rightarrow K_4[Fe(CN)_6]+2H_2O+2CO$
 - (b) potassium cyanide and ferrous sulphate solution. FeSO₄+2KCN→Fe(CN)₂+K₂SO₄ Fe(CN)₂+4KCN→K₄[Fe(CN)₆]
- Potassium Hydrogen Carbonate, KHCO₃. Precipitated from an aqueous solution of K₂CO₃ and CO₂. Used in foods, medicines
- Potassium Hydrogen Tartrate, Cream of Tartar, C₄H₅O₆K. Colourless salt, soluble in boiling water, occurs in grape juice, deposited as argol during fermentation. Used in baking powder.
- Potassium Hydroxide, KOH (Caustic potash). A white soild (M.P. 306°C) obtained by electrolysis of KCl solution. A strong base electrolyte in the Ni-Fe electric storage battery and in the molecules of water. Forms hydrates with 1, 1½, and 2
- Potassium Iodate, KIO₃. A white solid obtained from I₂ plus KOH solution. Used in volumetric analysis.
- Potassium Iodide, KI. A white ionic solid obtained from l_2 and hot KOH solution. Soluble in water and dissolved iodine to form polyiodide ions $[I_3]^-$, $[I_5]^-$, etc.

Potassium Nitrate, KNO3 (saltpetre, Nitre). A white solid, soluble in water, formed by fractional crystallization of sodium nitrate and potassium chloride solution. Occurs in nature as saltpetre. It is non-deliquescent and used in gun powder, fertilizers and in the laboratory preparation of HNO3.

Potassium Nitrite, KNO₂. A creamy deliquescent salt (M.P. 440°C) obtained from KNO₃ and lead. Used in diazotization.

Potassium Oxalate, C₂O₄K₂, H₂O. Used for removing ink stains and iron mould. Obtained from KOH or K2CO3 and oxalic

Potassium Perchlorate, KClO₃. Prepared by heating KClO₃ to

Potassium Permanganate, KMnO₄. A purple solid (soluble in water) obtained by oxidizing potassium manganate (VI) with Cl₂. Used as an oxidizing agent in volumetric analysis, as a bactericide and as a disnifectant. Acts as an oxidizing agent in acidic as well as alkaline solution (pH-dependent).

Potassium Sulphate K₂SO₄. A white solid prepared by the neutralization of either KOH or K₂CO₃ with dil H₂SO₄. Occurs in glaserite, schonite and syngenite. Used as a fertilizer (for to-

bacco plants).

Potassium Thiocyanate, KSCN. Obtained from KCN and S Used

in volumetric analysis.

A method in which the electrode dipped in the solution shows a rapid change in electromotive Potentiometric Titrations. force as the end point is reached. The end point is determined by noting the change in e.m.f. of the electrode during the titration. Used mainly for coloured solutions.

Potter's Clay. Describing a clay used for making pottery. Coloured clays are obtained by adding coloured oxides to

Powder Metallurgy. A process of sintering powder below the M.P. to produce solids. Used for alloy and some plastics.

Power Kerosine (TVO). Tractor vaporizing oil.

Pr. Praseodymium (also propyl).

Praseodymium, Pr. At. No. 59, At. Wt. 140 90, M.P. 931°C, B.P. 3212°C, D, 6.77. A lanthanide. Metal is used in thermoelectric material and glasses. Forms Pr3+ (pale green) compounds.

Precipitation. A process by which an insoluble compound is formed from its solution either by interaction of two salts, e.g.

KC!+AgNO₃→AgCl ↓ +NaNO₃ (in solution)

or by changing temperature which effects solubility.

Precipitation Hardening, Age Hardening. A process of hardening of some alloys which take place with time when a supersaturated solution tends to decompse with partial precipitation of the solute metal as an intermetallic compound.

Precipitation Indicator. An indicator which forms coloured precipitate at the end point, e.g. potassium chromate in estimation of ions by Ag⁺ ions (red ppt of Ag₂ CrO₄).

Predissociation. A term used in spectroscopy.

Prednisolone, C₂₁H₂₈O₅. White crystalline powder, obtained by the action of certain micro-organism on hydrocortisone. Used medicinally for treating asthama, rheumatic fever, etc.

Pregnane, 17β-ethylaetiocholane, C₂₁H₃₆. A basic skelton of biologically and clinically important steroids.

Pregnanediol 5 β -pregnane-3 α 20 α -diol, $C_{21}H_{36}O_2$. M.P. 238°C. Present in liver and is the chief urinary metabolite of it.

Prehnitene, 1,2,3,4, Tetramethyl benzene, C₆H₂ (CH₃)₄.

Prephenic Acid, C₁₀H₁₀O₆. An intermediate in the 'shikimic acid' pathway, of aromatic biosynthesis, arising by rearrangement of chorismic acid.

Primeverose, C₁₁H₂₀O₁₀. A disaccharide (glucose 6-β-D-xyloside), M.P. 208°C.

Pressure, P. Force acting on a surface per unit area:

The unit is Pascal (Pa).

Primary Standard. A substance that can be used directly for the

Primitive Lattice. A lattice having only one equivalent point in the unit cell.

Principal Quantum Number (n). Defines the number of orbit of an electron in an atom, e.g. K (n=1), L (n=2), M (n=3), and N (n=4).

Printing Ink. A sol in which a mixture of pigments (e.g. carbon black) dispersed in an oil of suitable viscosity.

Procaine, Diethylaminoethyl-P-Aminobenzoate. NH2 C6H4. COOC2 H₄. N (C₂H₅)₂. C₁₃H₂₀N₂O₂. A powerful local anaesthetic. Its hydrochloride has M.P. 156°C.

Procion Dyes. A type of azo dyestuffs which can form covalent

bonds to cellulose by reactive groups.

Producer Gas (air gas). A mixture of CO (25-30%), nitrogen (50-55%) and hydrogen (10-15%), produced by passing air with a little steam through a thick hed of white hot coke in a furnance (producer). Its calorific value (from coke) is 46,00 KJ/m³ and about 6400 KJ/m3 (from coal). Progesterone, C₂₁H₈₀O₂. A hormone of corpns luteum which

governs the development of uterus during pregnancy. Proguanil, C₁₁H₁₆ClN₅. Prepared as its hydrochloride (M.P. 248°C). Used for the treatment of malaria.

CI NH.C.NH.C.NH.CH

Prolactin. A hormone present in pitnitary gland. Proline, 2-Pyrrolidine Carboxylic Acid, C5H9NO2. Colourless crystalline solid (M.P. 22°C) Not exactly amino acid.

Promethazine, C₁₇H₂₀N₂S. A white powder (M.P. 223°C). A powerful antagonist of histamine. Used for the treatment of

Promethium. At. No. 61, At. Wt. 145, M.P. 1680°C, B.P. 2460°C, D 7.22. A radioactive element of the lanthanoid series, produced artificially by the fission of uranium. Forms a single series of typical lanthanide compounds in the +3 state. Used in luminescent paint for watch dials.

Promoter (activator). A substance that improves the efficiency of a

Proof Spirit. A solution of ethanol in water which at a temperature of 51°F (283.5K) weighs exactly 12/13 of an equal volume of distilled water. This also corresponds to an alcoholic solution of relative density 0.9198 at 15.6°C (288.6 K) and containing 57.1% v/v or 49.2 w/w of ethanol.

1.2-propadiene, allene, $CH_2=C=CH_2$, C_3H_4 . Colourless gas (Zn plus alcohol on 1,3-dibromopropane) which is isomerized to

propyne (methyl acetylene).

Propanal, Propionaldehyde, CH₃ CH₂ CHO. Colourless liquid, B.P. 48°C. Prepared by dehydrogenation of n-propanol over a 48°C. Frepared over a catalyst (Fischer-Tropsch) or oxo combination of C₂H₄, CO and

Propane, C₂H₈. A gaseous alkane obtained either from the gaseous fraction of crude oil or by the cracking of heavier fractions. Used as a fuel for heating and as a refrigerant. (M.P.—190°C, B.P.—44.5°C) B.P. -44.5°C).

Propanedioic Acid. See malonic acid.

1, 3-Propane dithiol, HS (CH₂)₂SH. Liquid (B.P. 170-171°C). Used in organic reactions for protecting a carbonyl group at a trimethylene dithioketal whilst transformations are affected at the other centres of the model. other centres of the molecule.

 $\left(> C \left\langle \frac{S - CH_2}{S - CH_2} \right\rangle CH_2 \right)$

Propanoic Acid, Propionic Acid, C₃H₆O₂, CH₃CH₂COOH. A colourless liquid carboxylic acid (M.P. 24°C, B.P. 97°C) occurring as a wood distillation of proposed and distillation product. Prepared by the oxidation of propanol and propenoic acid. Used for the manufacture of polymers.

1-Propanol, n-Propyl Alcohol, C₃H₈O, CH₃CH₂CH₂OH₄. Colourless volatile flammaki 100°C) volatile flammable liquid with a pleasant odour (B.P. 97°C) Occurs in fusel oil. Prepared by hydrogenation of propene oxide. Forms propanal and propionic acid on oxidation and propene when heated over alumina. Used as a solvent.

2-Propanol, Isopropyl Alcohol, C3H8O, CH3 CH2OH CH3. A colourless volatile flammable liquid (B.P. 82°C). Forms acctone (oxidation); amines, esters, glycerol etc. Used as a solvent for

several resins, aerosols, antifreezes.

Propanone, Acetone, Dimethyl Ketone, C_3H_6O , CH_3 C=O

A colourless liquid, B.P. 56°C and highly inflammable. Manufactured by the delay. factured by the dehydrogenation of 2-propanol over a copper catalyst at 500°C and 4 atm.

CH₃ CH OH CH₃ $\xrightarrow{\text{H}_2\text{O}}$ CH₃ CO CH₃ Acetone

Formed by the oxidation of cumene. Occurs in blood and urine. Forms diacetonamine with ammonia (boil), diacetone alcohol (condensation with NaOH) and phorone. Reduction of propanone produces propanol and pinacol. Forms crystalline derivative with phenylhydrazine, hydroxylamine and semicarbazide. Detected by the formation of iodoform (alcohol+12+ NaOH). Does not reduce Fehling's or Tollen's reagents. Used as a solvent and for the manufacture of methyl-methacrylate.

Propargyl Alcohol. See 2-propyn-1-01.

Propallants. Describing an explosive used to propel rocket, bullet, shot or shell. Also denotes liquefied gas in a pressurised aero-

Propenal, Acrolein, Acraldehyde, Vinyl Aldehyde, CH2 CH-CHO. Colourless volatile liquid with a typical odour,

B.P. 530°C. Highly poisonous and intensely irritating to eyes. Manufactured by the direct oxidation of propene or cross-condensation of ethanal with methanal. Prepared by dehydration of glycerol by heating with potassium hydrogen sulphate.

Converts to a white insoluble resin (disaeryl) when exposed tolight. Forms propenoic acid (on oxidation) and dibromide with

Br₂). Used to produce acrolein polymers.

Propene, Propylene, C₃H₆, CH₃CH=CH₂. A colourless gaseous alkene (B.P. - 48°C) obtained by cracking petroleum or by passing the vapour of 1-propanol over heated alumina. Forms cumene with benzene. Used for the manufacture of polymers,

propanone, etc.

Acrylonitrile, Vinyl Cyanide, CH2=CH-CN. Volatile liquid, B.P. 78°C. Manufacture by the catalytic de-Propenenitrile, hydration of ethylene cyanhydrin or by the addition of hydrogen cyanide to ethyne in the presence of CuCl. It is very active and forms polymers and copolymers which are used as synthetic fibres, nitrile rubbers and in thermoplastics. Used as an active fumigant against stored grain insects.

Propene Polymers, Polypropene, Polypropylene. An important class of polymers used as moulding resins and in extruded form,

Propenoic Acid, Acrylic Acid, Vinylformic Acid, CH2=CH.COOH. Colourless liquid, B.P. 141°C. Prepared by the oxidation of propenal with moist Ag₂O or treating β hydroxy propionitrile with H₂SO₄. Converted slowly into resin at ordinary temperature. Polymerized to several important polymers.

Propenol, Allyl Alcohol CH2=CHCH2OH. Colourless liquid, B.P. орепоі, Anyi Attonion glycerol and oxalic acid or manufactured by direct chlorination of propene at 500° C followed by hydrolysis of allyl chloride. Estimated as its bromine derivative. Oxidized to glycerol by KMnO₄. Forms propenal and propenoic

Propenyl Isothiocynate, Allyl Isothiocynate Mustard Oil, C4H5N5 CH₂=CHCH₂NCS. Colourless, intensively pungent liquid, B.P. 151°C. An important constituent of black mustard oil. B.P. 151 C. Prepared by reacting propenyl iodide in alcohol with potassium thiocynate. Used as a counter-irritant.

Propenylthiourea Allylthiourea, Thiosinamine, Rhodailin, C4H8N2S CH₂=CH.CH₂NHCSNH₂. Colourless solid, garlic like smell, M.P. 74°C. Manufactured from propenylisothiocynate and NH₃ in alcohol. Used as a chemical sensitizer for photographic

silver halide emulsions and medicinally for treating excessive

β-Propiolactone, $C_3H_4O_2$, $\left\{ \begin{array}{c} CH_2-CH_2 \\ | \\ O--C=O \end{array} \right\}$

Colourless liquid, BP. 162°C (decomposes). A highly reactive substance, prepared from methanol under controlled conditions (anhydrous ZnCl2). A potent carcinogen.

Propranolol, 1-(Isopropylamino)-3-(1-Naphthyloxy)-2-Proponol, C₁₆H₂₁NO₂. Prepared as its hydrochloride, M.P. 163°C. Used modicinally for treating hypertension.

Propyl. The C₃H₇-group. Propyl Alcohol. See propanol.

Propylene Chlorohydrins, C₃H₇ClO. a Propylene chlorohydrin, 1-chloro-2-hydroxy propylene Cliffold a Propylene chlorohydrin, 1chloro-2-hydroxy propane, CH2Cl.CHOH, CH3, B.P. 127°C. B.P. 134°C. chlorohydrin, 2-chloropropanol, CH3CHCICH2OH.

Both are colourless liquids obtained by treating dilute solutions
Na HCO and propene-oxide E of HO Cl and propene-oxide. From 1, 2-dihydroxypropane with Propylene Glycol. See 1, 2 Dihydroxy propane.

Propylene Oxide, 1, 2-epoxypropane, CH₃CH.CH₂

Colourless liquid, B.P. 34°C. Prepared by heating chlorohydrin produce polystrom NaOH. Recember by heating chlorohydrin Used to with solid CaO or NaOH. Resembles ethylene oxide. Used to produce polyglycols and as a solvent for nitrocellulose.

Propyne, Allylene, Methylethyne, CH₃C CH. An ethyne, B.P. propane.

2-Propyn-1-01, Propiolic Alcohol, CH C-CH2OH. B.P. 112°C. Prostaglandins. A group of related unsaturated hydroxylated fatty

Occurs in mammalian organs, tissues and secretions. Inhibits arachiaggregation of blood platelets in body. For example, arachiaggregation of process of body. For exam donic acid, a biological precursor to prostaglandins.

Prosthetic Group. The non-protein group of a conjugated protein.

At No. 91 At W. ntactinium, Pa. St. 15.4, At Wt. 231.036, M.P. 1565°C (half-life 3248) At No. 91, At Wt. 231'036, M.P. 1565°C, D toxic radio active element of the actinoid series of metals. Occurs in U ores as a radioactive decay product of Actinium. Originally separated from pitchblende residue by co-precipitation with ZrO_2 followed by removal of Zr as $ZrOCl_2$ and of Ta as the soluble peroxide. The silvery metal shows oxidation states of +5 and +4. Forms PaH_5 (colourless), $PaCl_5$ (yellow) and PaH_3 (with H_2).

Protecting Group. A group which when added to one functional group in a molecule before chemical treatment of a different functional group, prevents unwanted reaction at the site. Used very often in synthesising peptide from aminoacids.

Protective Colloids. Hydrophilic colloids which when added in small quantities prevent hydrophobic sols from coagulating influence of electrolytes. For example, gelatin, starch and casein.

Preteins. A naturally occurring compound present in living matter consisting of amino-acids linked into long chains. They are chief hydrogen constituents of living organisms containing about 50%C, 25% oxygen, 15% nitrogen, about 7% hydrogen and some S. Proteins are precipitated by alcohol, propanone and by strong solutions of salts. Important proteins are classified as follows:

Simple Proteins, e.g. albumins (ovalbumins from eggs), globulins, (myosin, fibrinogen), protamines (soluble proteins), low Mt. wt., histones (basic, act as specific inhibitor of nuclear genes) prolamines (present in seeds of cereals, insoluble in water), conjugated proteins (contain prosthetic group), nucleoproteins (contain prosthetic group as nucleic acid, linked through salt linkages with protamines or histones).

Lipoproteins. (a-lipoprotein of serum contains glyceride, phosphatide and cholesterol to about 30—40% of the total complex; B-lipoprotein contains about 75% of phospatide and cholesterol), glycoproteins or mucoproteins (protein compounds with hydrocarbons, have covalent or salt like linkage).

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where R and R' are amino acid residues. Proteins may have more than one peptide chain. In the fibrous proteins the polypeptide chain is in the form of a regular helix. The structure of proteins is determined by X-ray analysis. They have M. wts-varying from 60000000 to 50000.

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Protocatechuic Acid, 3-4-Dihydroxybenzoic Acid, M.P. 199°C. Occurs in onion and a constituent of a group of tannins. A product of decomposition of resins (alkaline).

Proton. An elementry particle and one of the units from which all forms of matter are built. It is identical to the nucleus of the hydrogen atom having a mass of 1 atomic unit and a unit positive change equivalent to the negative change on the electron. Proton is a nucleon and a proton is 1840 times the

Proton Number (atomic number), Z. The number of protons in the nucleus of an atom. It determines the chemical properties

Protonation. Describing the addition of a proton, e.g. protonation of H₂O by HCl to produce (H₃O)+Cl⁻ ion.

Protonic Acids. Molecules which produce H+ ions on dissociation,

Protophilic. Proton loving.

Prussian Blue. KFe Fe(N)6.

Prussic Acid, HCN. A name for HCN.

Pseudobrookite Structure. Describing structures of the type A₂BO₅ materials, where A has values between 4 and 6 co-ordination and B is octahedrally co-ordinated e.g., Fe₂TiO₅.

Pseutocumene, 1, 2, 4-Trimethylbenzene, C₆H₃Me₂. A liquid used

Pseudo-First Order (reactions). Chemical reactions which look like of second order but in practice they are first order reactions.

Pseudohalogens. Compounds which resemble halogens in their properties, e.g. (CN)₂, (SCN)₂, (Se CN)₂, etc.

Pseudomorphic. Describing species of same crystalline form.

Psychotomimetic Drugs, Psychopharmacological Agents. Drugs which produce changes in perception, thought and mood without disturbing the autonomic nervous system, e.g., lysergic acid diethyl amide (LSD-25) and mescaline.

Pt. Platinum.

PTA. p-Terephthalic acid.

Pterins. Derivatives of pteridine

$$\begin{bmatrix}
8 & 1 \\
N & N \\
7 & & 2 \\
6 & & 3 \\
N & N \\
5 & 4
\end{bmatrix}$$

Pu. Plutonium.

Pulp. A fibrous cellulose material used in paper, rayon, cellulose production. It is made from wood, cotton, linean, straw, etc., by mechanical or chemical treatment (e.g. Na₂S or Na OH, etc.).

Pump. A mechanical device for moving a fluid.

Purine, C₅H₄N₄. A crystalline solid, M.P. 217°C.



Purple of Cassius. Colloidal gold.

Putrescine, Tetramethylenediamine, H₂N (CH₂)₄ NH₂.M.P. 27-28°C.

Present in ergot, urine, etc.

Putty Powder. See tin oxide.

PVC. Polyvinyl chloride.

Pyknometer. An instrument for determining density.

Pyranose. A sugar with a six membered ring (five C atoms and one oxygen atom), e.g. glucose is a 1,5-glucopyranose with a primary alcohol group CH2 OH as a side chain.



Pyrazine. 1,4-Diazine.



Pyrazole, 1,2-Diazole, C₃H₄N₂. Colourless substance (M.P. 70°C), obtained by passing ethyne through a cold etheral solution of obtained by passing children of diazomethane. Undergoes electrophilic substitution in position





Obtained by complete reduction

of pyrazole.

Obtained by partial reduction of

pyrazole.

Pyrazolones, Pyrazolinones. Prepared from acetoacetic ester and phenylhydrazines (e.g. Fast yellow G). Pyrene. A colourless tetracyclic benzenoid hydrocarbon, M.P.

150-151°C.

Pyrethrolone. A ketonic alcohol which forms pyrethrins (ester). Pyrethrum. A powerful but, non-persistent insecticide with rapid knockdown effect. A mixture of dried flowers of Chrysanthemum cinerariaefolium.

Pyrex. A trade name for borosilicate glass having high percentage of SiO, and some P. Al missilicate glass having high percentage of SiO₂ and some B, Al., alkalis. It is heat resistant, resists attack by strong acids and alkalis and has high mechanical strength.



Pyridine, C₆H₅N, 5/3 6/2. B.P. 115'3°C. A heterocyclic nitro-

gen compound with the electrons in the C-C, m bonds and the lone pair of the nitrogen delocalized over the ring of atoms. Extracted from coal tar and manufactured from ethyne and ammonia. Used as a solvent in plastic industry and in the preparation of nicotinic acid.

Pyridinium. The ion



Pyridoxal Phosphate. A prosthetic group of transmination i.e. the interconversions of amino-acids and their corresponding a-keto acid.

Pyridoxine, Vitamin-Be, 2-Methyl-3-hydroxy-4, 5-hydroxy-methyl

Pyridine, C₈H₁₁NO₃. Colourless crystalline, M.P. 160°C. Present in rice, husks, maize, wheat germ, yeast and other sources of vitamin B. Supplemented by thiamine and riboflavin. Causes anaemia if absent from diet.

Pyrimidine, C₄H₄N₂. Crystalline, M.P. 20-22°C, B.P. 124°C. Obtained from barbituric acid.

Pyrites, FeS2. A mineral used for production of H2SO4.

Pyro Acids. Acids or anions obtained from hypothetical acids having two oxygroups with a bridging (μ) oxygen, e.g. $(O_3 \text{ SO } SO_3)^{2-}$.

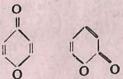
Pyrochlore, Pyrochloride, NaCaNb₂O₆F. An important mineral of Nb.

Pyrogallol, 1,2,3-Trihydroxybenzene, C₆H₆O₃. White needles, M.P. 132°C, B.P. 210°C. Prepared by digesting gallic acid with water at 200°C. Absorbs (in alkaline solution) O₂ from the air and turns dark brown in colour. Used as a photographic developer.

Pyroligneons Acid. Obtained from wood distillation.

Pyrolusite, β-MnO₂. An ore of Mn.

Pyrones. Compounds having the ring and occur in nature.



Pyrophoric Metals. Metals which when produced in a porous condition or finely divided state by reduction at low temperature are inflammable and in appreciable activity, e.g. Ce-Fe alloy used in lighters.

Pyrophosphates. Salts having [P2 O7]4- ions.

Pyrosols. Prepared by electrolysis of fused salts or by dissolving the metal directly in the fused salts, e.g. Zn in fused Zn Cl₂.

Pyrosulphuric Acid, H₂S₂O₇. Acid containing [S₂ O₇]²⁻ ion. Present in fuming sulphuric acid.

Pyrosulphurous Acid, H₂S₂O₆. Formed from sulphites and sulphur dioxide. It contains [S₂O₅]² ion.

Pyrotartaric Acid, Methyl Succinic Acid, C5H8O4.

CH₃ CH COOH

CH₂ COOH

CH₂ COOH

lation of tartaric acid

Pyrotechnics. Describing combustible mixtures containing an oxidant such as nitrate or chlorate and combustible substance (e.g., ing and smoke-producing composition.

Pyroxenes. A class of silicate minerals containing (Si O₃)₄²ⁿ⁻, e.g.,

Pyrrole, C₄H₅N, . A heterocylic colourless oil, B.P.

130°C. It is aromatic containing π (delocalized) electrons. Prepared from butyne-1, 4-diol and ammonia in the presence of ammonia catalyst.

Pyrollidine Tetrahydropyrrole, 3 4 C₄H₉O. Ammonia

like liquid, B.P. 88-89°C. H Occurs in tobacco leaves. H strong base.

Pyruvic Acid, 2 Oxopropanoic Acid, CH₃COCOOH. A ketoacid and a colourless liquid, M.P. 130°C, B.P. 65°C/10 mm. Prepared by distilling tartaric acid with KHSO₄. Reduced to (±) lactic acid.

Q

Quadrivalent. Showing a valency of four.

Quadrupole Moment. The coupling of the nuclear quadrupole with the asymmetric electron field in compounds. It provides informolecule or ion.

Quadrupole Point. The condition of T and P at which four phases of a two-component system are in equilibrium.

Qualitative Analysis. Analysis for the identification of components of a mixture.

- Quantitative Analysis. A method for the estimation of constituents of a mixture or species.
- Quantized. Describing a physical quantity that can only take certain discrete values, and not a continuous range of values. For example, in atom and molecules, the electrons have quantized values of spin angular momentum and orbital angular momentum. The energy levels E₁, E₂... E₄ of an atom are quantized.
- Quantum. According to Planck's quantum theory the transition of energy can only occur in definite parcels, or quanta, and not continuously as a stream of fluid. The quantum of electromagnetic radiation is 'photon'. For radiations of frequency v, the unit of energy, the quantum, is equal to hv where hv is Planck's constant.
- Quantum Efficiency. It is the number of molecules actually decomposed for each quantum of radiation absorbed.
- Quantum Electrodynamics. Describing the use of quantum mechanics to explain the interaction of particles and electromagnetic radiation.
- Quantum Number. The electrons moving around the atom or atoms in a molecule possess energy due to rotation, vibration or spin. According to the quantum theory the energy possesed by such a molecule is quantized. Each such value, for a particular form of energy (e.g., rotational) is a multiple-which is always a small whole number, or sometimes one-half of a unit of energy, the quantum, which is characteristic of a specific type of energy under consideration. This multiple which defines the energy of a particle, is termed as quantum number.
- Quantum Mechanics. According to Broglie, particles can have wave like properties-this type of quantum mechanics is known as wave mechanics.
- Quantum States. States of an atom, electron, particle, etc. Characterised by a unique set of quantum numbers. For example, hydrogen atom in ground state has only one electron in K (n=1) shell and is specified by four quantum numbers, e.g., n=1, l=0, $m=\pm\frac{1}{2}$.
- Quantum Theory. A mathematical theory introduced by Max Planck and which is based on the view that energy (or some other physical quantities) can be changed only in certain discrete amounts for a given system.
- Quark. A fundamental sub-atomic particle.
- Quartering. A common method of obtaining a small representative sample of a solid material.
- Quartet. A group of four closely spaced lines in a spectrum.
- Quartz. A naturally occurring crystalline form of silica (SiO₂). It is insoluble in all acids except HF. Hardness, 7, Sp. gr. 2.65 M.P. 1600°C-1700°C.

Quaternary Salts. The products of reaction of e.g. teritary amines or phosphines with alkali halides. Ionic Salts. These salts are used in fabric softening (cations react with fabrics), as germici-

Quenching. A process for altering the mechanical properties of metals, e.g., hardness. The hot metal is quickly dipped in an oil, water or brine followed by rapid cooling. Also applied to

Quercitrin, C21H20O4. A glycoside widely distributed in plants. M.P. 169°C. Used as a natural dyestuff.

Quick Lime. CaO.

Quick Silver. An old name for mercury.

Quinaldine, 2-Methylquinoline, C₁₀H₉N. Colourless oil, 246-247°C. Used for the preparation of photosensitizing dyes.

Quinhydrone, C₁₂H₁₀O₄. A stable compound constituted by quinone and hydroquinone molecules linked together by hydrogenbond. M.P. 171°C. Used in the quinhydrone electrode.

Quinhydrone Electrode. An electrode used for measuring the hydrogen-ion concentration in a solution, or for measuring the electrode potential of a substance. The reaction of the electrode

 $C_6H_4O_2+2H^+ \rightleftharpoons C_6H_6O_2-2e^-$

Quinic Acid, C₆H₇ (OH)₄ COOH. A carboxylic acid of a tetra-hydroxycyclohexane. Present in cinchona bark, coffee beans,

Quinidine, C₂₀H₂₄O₂N₂, 2H₂O. A (+) isomer of quinine. Occurs in cinchona bark and a by-product in quinine preparation.

Quinine, C₂₀H₂₄O₂₄N₂, 3H₂O. A white crystalline powder, M.P. 57° and 177° (anhydrons). Insoluble in water, partly soluble in alcohol. A principal alkaloid of cinchona bark obtained by mixing the bark with lime followed by extraction with alcohol of

by boiling the bark with acid followed by precipitation with ammonia. Purified via its sulphate. Also synthesized from 7-hydroxyisoquinoline.

Quanine and its salts are used for treatment of malaria (destroys malaria parasites). Retards enzyme action. An anti-

pyretic used to treat cold, influenza, etc.

Quinitol. 1, 4-cyclohexanediol.

Quinizarin, 1, 4-dihydroxy-9, 10-anthraquinone, C14H8O4. M.P. 195°C. Used for the preparation of several dyestuffs.

Quinol. Hydroquinone.

Colourless oily liquid B.P. Quinoline, C9H7N,

238°C. Hygroscopic; occurs in high boiling fraction of coaltar. Basic, forms stable salts with mineral acids and ammonium compounds. Prepared by Skraup's method. Used in the manufacture of pharmaceuticals.

Quinolinol. 8-Hydroxyquinoline.

Quinone. Benzoquinone.

Quinones. Highly coloured substances derived by replacing the >CH₂ groups in a di-hydroaromatic system by >C=O. Used as dehydrogenating agents.

Quintet. A group of five closely spaced lines in a spectrum.

Quinuclidine, C₇H₁₈N. A strong, hindered base (PKb=11), M.P. 158-159°C. Used for metalation.

Racah Parameters. The parameters used to express quantitatively the inter-electronic repulsion between the various energy levels of an atom. Expressed as B and C.

Racemic Mixture. An equimolecular mixture of the dextro (+) and

laevorotatory (-) optically active compound.

Racemization. The conversion of an optically active isomer into an equal mixture of isomers which are optically inactive. Racemization is brought about by heat and the action of acids.

Radian (rad). The SI unit of plane angle; 2π radians in one complete revolution (360°).

Radiation. The emission of energy from a source either as waves (e.g., light, sound etc.) or as moving particle (e.g., β-rays or

Radical. Species having one or more free valencies, e.g., free

Covalent and ionic radii of elements.

Radioactive. Describing an element or nuclide that shows natural radioactivity.

Radioactive Decay Series. Describing the series of isotopes into which a radioactive nucleus is successively transformed.

Radioactivity. Becquerel (1896) found that certain unstable nuclides (e.g., U) disintegrate with emission of radiation such as a-, β-and γ-. The emission of an α-particle results in decrease in mass of the nucleus by four units; β-particle emission corresponds to an increase in atomic number of the nucleus by one unit and the emission of \(\gamma\)-rays results in loss of energy of the nucleus (other accompanied by x-rays). All elements having At. No. more than 83 show radioactive decay. Radioactive decay is always spontaneous. Position emission corresponds to a decrease by one unit in atomic number, whereas, K capture results in the incorporation of one of extranuclear electrons and At. No. is decreased by one unit.

Radioactivity, Artificial. It was shown that there are certain elements which undergo nuclear reaction when bombarded by a-rays. The elements also undergo a nuclear reaction when bombarded with protons, deuterons or neutrons. All these resulting atoms may be stable or metastable and show radioacti-This phenomena is known as artificial or reduced radioactivity. Some of the artificially radioactive substance give out positrons or positive electrons.

The reaction between the alumunium nuclei and a-particles is given as follows:

 $^{27}_{13}\text{Al} + ^{4}_{2}\text{He} \rightarrow ^{30}_{15}\text{P} + ^{1}_{0}n.$

where ${}^{1}_{0}n$ is a neutron. The new phosphorus nucleus has characteristics of radio. characteristics of radioactivity and disintegrates into atoms of a silicon isotope and positive electrons:

 $^{30}_{15}P \rightarrow ^{30}_{14}Si + ^{1}_{0}e$

This process produces the observed artificial radioactivity. Neutrons are also quite effective in producing artificial radioactivity. A large number of the control of the c vity. A large number of radioactive isotopes are radio-iodine radio-sodium, radio-silver, etc.

Radiocarbon Dating. All living things contain radioactive 14C in a uniform quantity. This 14C is produced constantly in atmos phere by the action of cosmic rays. When the thing is dead, the exchange of non-radioactive carbon with the atmospheric 14C is also stopped. So by determining the amount of ¹⁴C in the host (dead sample), the age of it may be determined. The half-life of 14C is approximately 5570 years.

Radio Waves. A form of electromagnetic radiation with wave lengths greater than a few millimetres.

Radium, Ra. At. No. 88, At. wt. 226.025, M.P. 700°C, B.P. 1140°C. D, 5. A white radioactive luminescent metallic element of the alkaline-earth group. Has various isotopes of various decay series. 226Ra (half life 1600 years) is least active. Isolated from U ores, such as the oxide pitchblende and carnallite by electrolysis using a Hg electrode. Electronic configuration 7s². Ra Cl₂, Ra F₂ and Ra Br₂ are known. Used in self-luminous paints, as a neutron source and in radiotherapy.

Radon, Rn. At. No. 86, At. Wt. 222, M.P. -71°C. B.P. -61'8°C. A colourless monoatomic radioactive element of the rare-gas group. A product of radioactive decay of the heavy elements. Has 19 short-lived isotopes and the most stable is 222Rn. An appreciable hazard in \cup mines. Electronic configuration $6s^26p^6$. Disintegrates into an isotope of polonium. Used in radio-

Raffinose, Melitose, C18H32O16. A prismatic pentahydrate (M.P. 80°C) trisaccharide containing galactose glucose and fructose

Present in cotton-seed meal and sugar beet.

Raman Effect. The light scattered by molecules of a substance when analysed spectroscopy has lines of frequency v,

v=vo+vi

and vo is the frequency of light (original) and vo is the vibrational frequency of molecules of the substance. This spectrum is called Raman spectrum and corresponds to the rotational and vibrational changes in the molecule.

Ramsay-Shields Equation. An equation

$$\begin{array}{c}
\text{Say-Shields Equation.} & \text{The options} \\
\gamma \left[\begin{array}{c} M \\ D \end{array} \right]^{2/3} = K \left(T_0 - T - 6 \right)
\end{array}$$

where γ is the surface tension of the liquid, M is its molecular weight, D its density, To is the critical temperature, T is the temperature at which molecular surface energy is measured and K is Ramsay-Shields constant (approx 2.12 for undissociated

Raney Nickel. A catalytic form of nickel obtained by treating Ni-Al alloy with caustic soda (a spongy mass, becomes pyrophoric when dry). Used for catalytic hydrogenation reactions.

Raoult's Law. The vapour pressure of a solvent is lowered down by the addition of a solute (non-volatile) and the relative lowering of the vapour pressure is directly proportional to the mole

fraction of the solute in the solution $\left(\frac{p-p_s}{p} = \frac{n}{n+N} = X \text{ solute}\right)$

where p=vapour pressure of solvent, p=vapour pressure of solution, and n and N are the number of moles of solute and solvent respectively. It is strictly followed by an ideal solution.

Rare Earths. A term applied strictly to oxides of the lanthanide elements but often applied to the elements itself.

Rare Gases. The noble gases, e.g. He, Ar, etc.

Raschig Process. See hydrazine.

Rast's Method. A method of determining molecular weights by measuring depression in freezing point of a solvent by the addition of the solute whose molecular weight is to be determined.

Rational Indices, Law of. The intercepts of the faces of a crystal upon the various axes of a crystal bear a simple ratio to each other.

Ratio of Specific Heats. A characteristic of the number of atoms in a molecule of a gas (e.g., monoatomic, diatomic, etc.) and is Cp/Cv, where Cp and Cv are the specific heats of gas at constant volume respectively. (For monoatomic gas 1.67, diatomic gas 1'40 and triatomic gas 9'33).

Rayon. An artifical fibre prepared from wood pulp. Viscose rayon is prepared by dissolving cellulose in NaOH and CS2 to form xanthated cellulose. The solution is forced through a fine nozzle into an acid bath, which regenerates the fibres.

Rb. Rubidium.

Re. Rhenium.

Reactant. A substance taking part in a chemical reaction.

Reaction Rate. The rate of a chemical reaction expressed either as the rate of disapperance of reaction(s) or rate of formation of product(s).

Reactive Dyes. Dyes which can react directly with fibres, e.g., acrylamide dves.

The various environments (e.g. tanks, towers, tubes, etc.) in which the chemical reactions are carried out.

Reagent. A term commonly used for laboratory substances (chemical), e.g. NaOH, HCl, etc.

Realgar. The mineral As₄S₄.

Rearrangement. A process in which the groups of a compound rearrange themselves to form a new compound.

Reciprocal Proportional, Law of. The weights of two or more substances which separately react chemically with identical weights of another substance are also the weights which react with each other or are simple multiples of them e.g., 23 gm of Na react 1 gm of H₂ to produce NaH; 23 gm of Na react with 35.5 gm of Cl₂ to form NaCl; 35.5 gm of Cl₂ react with 1 gm of H₂ to form HCl. (Not a general law).

Recrystallization. A process for the removal of impurities from compounds (crystals).

Rectification. A process of separating a mixture into its components by fractional distillation. Used in petroleum industry.

- Reactified Spirit. A solution of ethanol in water containing 90% v/v ethanol. A 57.8° over proof (OP).
- Rectifying Columns (fractionating columns). A long vertical cylinder containing either a series of plates, or a packing material (of large specific area) and a condenser at the top and a vapourizing unit (reboiler) at the bottom.
- Recuperators, Regenerators. Heat exchangers used in furnance operations.

Red Lead. Pb3O4.

Redox. An abbreviation for reduction-oxidation process. Reduction and oxidation reaction go side by side.

Redox Catalyst. Describing a free radical catalyst which combines with a reducing ion or salt which increases the rate of free radical production. Used in polymerization reactions.

Redox Indicator. A substance which operates on a definite and narrow potential range and undergoes a definite colour change during its reversible oxidation or reduction, e.g. methylene blue (colourless-reduced, blue-oxidized).

Reduced Mass. For particle in harmonic motion, the reduced mass (µ) is given by

 $\mu = \left(\frac{mM}{m+M}\right)$

where m and M are the masses of the particle, respectively.

Reducing Agents. Substances which can bring about reduction and themselves get oxidized, e.g. Sn2+, SO2, hydroxylamine etc.

Reduction. A chemical process involving a decrease in the proportion of electronegative substituents, e.g. SnCl₄-SnCl₂ or the charge on an ion is lowered or the oxidation number is lowered, e.g. KMn7+O4 to Mn2+ or Fe3+ to Fe2+, etc.

Reductone, Glucic Acid, C3H4O3. Solid, M.P. 140° (decomp.). A reducing agent (alkali-hexoses).

Reductores. Enediols stabilized by conjugation to a cabohyl or

Reference Electrode. Standard electrode used in electrochemistry, e.g., calomel electrode, silver-silver chloride (Ag-AgCl) electrode.

Refining. A process of removing impurities from a substance or of extracting a substance from a mixture.

- Refining Petroleum. A process used for converting crude oil into a wide range of petroleum products including fuels, lubricants,
- Reflux. A process used in organic chemistry in which a liquid from partial condensation of vapours returned to the top of a fractionating coloumn and allowed to run back the column counter current to the ascending vapour.

- Reforming. The cyclization of straight chain hydrocarbons from crude oil by heating under pressure in the presence of Pt or Al, e.g. the formation of methyl benzene from heptane
- Reformatski Reaction. A reaction in which carbonyl compounds react with α-bromo-fatty acid esters in the presence of Zn-powder to form β-hydroxy esters (α-chloroesters react in presence of Cu) followed by dehydration to produce α-, β unsaturated esters.
- Refrigerants. Materials used for cooling, e.g., liquid O₂ (-183°C), solid CO₂ (-78.5°C), etc.
- Refrigeration. A process for achieving low temperatures, i.e. achieving refrigeration, by vapour compression. For example, ammonia and freons (chlorofluorocarbons) vapours are used as-refrigerants.
- Regioselectivity. Tendency of reactions for proceeding in two or more ways to proceed in one manner only.
- Regnault's Method. A method used for determining the density of gases.
- Regular System. The cubic system.
- Reinforced Plastic. Plastics that have the thermosetting resin built up by heat and pressure.
- Relative Atomic Mass, Ar. The ratio of the average mass per atom of the naturally occurring element to $\frac{1}{12}$ of the mass of ¹²C. Formerly known as atomic weight.
- Relative Density, D or d. The ratio of the given substance to the density of some reference substance, e.g., for liquids the ratio to density of water at 4°C or for gases with respect to air at STP.
- Relaxation. The process of loss of energy from an excited state to the ground state or to another excited state.
- Rennin. A milk-clotting enzyme found in the fourth stomach of ruminants. Used in cheesemakings (rennet).
- Reserve Acidify and Alkalinity. A buffer solution, e.g., an equimolar mixture of acetic acid and sodium acetate.
- Resins. Materials with high molecular weights (controlled) and which soften at high temperatures.
- Resolution, (of racemics). A process for separating a racemate into its two enantiomorphic forms, i.e. (+)-and (-) forms.
- Resonance. The bonding in the molecule can be considered as a hybrid of two or more conventional forms of the molecules, called resonance forms or canonical forms. In optics it refers to

- the absorption of radiation (visible, IR etc.) by a system which it is capable of emitting. A term also applied to valency electrons in a molecule.
- Resonance, Ionization Spectroscopy. A technique applied to a single distinct atom by using a laser to ionize that atom selectively.
- Resorcinol, 1,3-Dihydroxybenzene, C₆H₆O₂. Needles, M.P. 110°C, B.P. 276°C. Prepared by fusing m-benzene disulphonic acid with caustic soda. Used for the manufacture of resins, plasticizers, etc.
- Respi atory Pigments. A class of pigments which act as oxygen carriers in living organisms, e.g., haemoglobin.
- Retene, C₁₈H₁₈. Yellow crystals, M.P. 101°C, B.P. 390°C. Occurs in tar-oils from pine wood.
- Retinene, Vitamin-A Aldehyde, C20H28O. Orange, M.P. 61-64°C Obtained from Vitamin-A.
- Retort. A laboratory apparatus used for distillation.
- Reverberatory Furance. A furnace for smelting metals.
- Reverse Osmosis. A process of separating solute from a solution by making the solvent to flow through a membrane at a pressure higher than the normal osmotic pressure. Used for concentration and dehydration of food products (e.g., milk and juices), for treatment of recycle water in chemical plants, etc.
- Reversible Process. A system in equilibrium can be made to act reversibly if it returned to the original starting position (or viceversa) by an infinitely change in one of the factors controlling the position of equilibrium (pressure, temperature, concentration), E.g. in reaction such as

 $2NH_3(g) \rightleftharpoons N_2(g) + 3H_2(g)$

which is at equilibrium, a slight decrease in pressure will shift the equilibrium towards right side, i.e., decomposition of ammonia.

Rf, Rs, Rr, Value. A measure of the relative distance travelled by a sample in chromatography and is the ratio of the distance travelled by the solute to the distance travelled by the mobile phase.

Rh. Rhodium.

rH. A concept used in redox reactions and is given by the expression: $rH = -\log p$

where p is the partial pressure of hydrogen in equilibrium with a system.

L-Rhamnose, C₆H₁₂O₅. Crystalises as 1H₂O (M.P. 94°C), anhydrous M.P. 122°C. A constituent of many glycosides present in flavanol derivatives.

Rhenium, Re. At. No. 75, At. wt. 186.2, M.P. 3180°C, B.P. 5627°C, D 20.53. A transition metal of Group VII with electronic configuration 5d. Shows oxidation states of +7 to -1. Obtained by reducing Re compounds with H₂. Used as an additive in W-and Mo-based alloys and in determining the age of universe. (Half life 7 × 10.10 years).

Rheopexy. A phenomenon in which the gelation of some thixotropic sols is accelerated by light mechanical agitation.

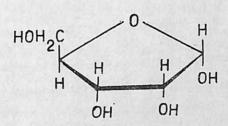
Rhodamines. Dyestuffs formed by Friedel-Crafts reactions in which phthalic anhydride and aminophenols are used.

Rhodium, Rh. At. No. 45, At. wt. 102'9, M.P. 1966°C, B.P. 3727°C, D 12'4. A platinum group transition metal with electronic configuration $4d^8 5s^1$. Shows oxidation states from +6 to -1. Difficult to work and highly resistant to corrosion. Obtained by extraction of ore with aqua-regia, fused KHSO₃, extraction with water to form Rh₂ O₃. Dissolve Rh₂ O₃ in HCl followed by reduction with H₂. Used as an alloying agent.

Rhodospin. An important photosensitive pigment of eye containing protein opsin and neoretinene-b.

Riboflavin, C₁₇H₂₀N₄O₆. M.P. 271°C. A part of the original vitamin B₂ complex which stimulates the growth of rats. A percursor of flavoproteins. Occurs in nature (e.g. liver milk, etc.).

D-Ribose, C₅H₁₀O₅. M.P. 87°C. A sugar of ribonucleic acid.



Ribosomes. Cell constituents (small, granular) of the endoplasmic reticulum. Have M. wt. of about 2.5 M. (for each ribosome) and containing several thousand in one cell.

Ribulose, C5H10O5. A key intermediate in the photosynthetic fixation of atmospheric CO.

Ricin. A protein of the albumin group. Highly poisonous, present in seeds of castor bean.

Ricinoleic Acid, C₁₈H₃₄O₃. M.P. 4-5°C. Cis-12-hydroxy-9-octadecenoic acid. Comprises about 85% of the acids of castor oil. Forms salts (ricimoleates).

Ring. A closed loop of atoms in molecules, e.g., in benzene

Ring Closure. Formation of ring from an open chain in a molecule.

Rinmann's Green, ZnCo2O4. Obtained by placing cobalt nitrate solution on Zn O and the mixture heated to redness. (A test for Zn).

Rn. Radon.

RNA. See nucleic acid.

Rochelle Salt, C₄H₄O₆ KNa, 4H₂O. Sodium potassium tartrate.

Rock Crystals, SiO2.

Rock Salt, NaCl. A transparent naturally occurring mineral form of NaCl.

Rodinol. See aminophenols.

Rontgen R. A unit of radiation and a measurement of ionizing. radiation (X-rays and Y-radiation). One rontgen induces 2.08 x 10° pairs of gaseous ions in air under standard condition.

Rontgen Rays. Discovered by Rontgen (X-rays).

Rose's Metal. A fusible alloy containing Bi (50%), Pb (25-28%) and Sn. Used in fire-protection devices.

Rosins. A solid resin used in the preparation of lacquers, plasticizers and flotation agents. Occurs in oil from pine trees.

Rotamers. Isomers obtained by restricted rotation.

Rotary Dryers. Devices used in industry for drying, mixing and sintering of solids.

Rotational Spectrum. The rotational energy of a molecule increases by absorbing energy and the emission of energy takes place in the transition from higher rotational energy state to lower rotational energy fals. The energy E emitted being given by E=hv and the lines in the molecular spectrum corresponding to such transition are known as rotational bands which of the molecule).

Rotatory Power Specific. The characteristic rotatory power of a pure liquid is expressed as

 $[a]^{t_{D}} = (a/1d)$

where $[a]^t = the$ specific rotatory power at temperature t for the sodium D line, a = the rotation of the plane of polarized light produced by a liquid of length l (decimeters) and density d. or tion of solute.

Rouge, Fe₂O₃. A finely divided form of Fe₂O₈ used for polishing. Royal Jelly. A nutrient of queen bee larvae.

Ru. Ruthenium.

Rubber. A natural or synthetic elastic polymeric material with high molecular weight. Natural rubber (obtained from the free 3-diene (isoprene). Synthetic rubbers are synthetic isoprene and butyl rubber. Vulcanized rubber is obtained by adding a reinforcing filler (e.g., carbon black).

Rubbone. A from of oxidized rubber.

Rubidium Rb. At. No. 37, At. wt. 85'468, MP. 38'89°C, B.P. 688°C. D 1'35 (solid) 1'47 (liquid). A soft silvery highly reacgroup element occurring in lithium mineral lepidolite. A first comprising of two isotopes (87Rb) and used in vaccum tubes, photocells and in ion-propulsion rocket motors.

Russel-Saunders Coupling, LS Coupling. A system in which electron spins are assumed to couple only with electron spins and orbital momentum to couple only with orbital momentum. Used to describe the ions of first row of the transition elements.

Rusting. The corrosion of iron in air to form a flaky layer of hydrated (III) iron oxide. Accelerated by SO₂ or H₂SO₄. Prevented by coating (Cr), painting, etc.

Ruthenium, Ru. At. No. 44, At. wt. $101^{\circ}07$, M.P. 2310° C, D $12^{\circ}45$. A transition metal with electronic configuration $4d^7 5s^1$. Shows oxidation state from +8 to -2. Occurs naturally with Pt and also forms alloys with Pt (Used in electrical contacts). A grey white metal obtained from its compounds by H_2 reduction. Used as a hardening agent for Pd and Pt.

Rutherfordium. Element 104.

Rutile Mineral TiO₂. A reddish brown mineral used as a source of T₄ in ceramics.

Rutin, C₂₇H₃₀O₁₆, 3H₂O. Obtained from buck wheat. Used in hypertension treatment.

Rydberg Constant. See Balmer series.

S

S. Sulphur.

Sabinene, C₁₀H₁₆. A dicyclic monoterpene colourless oil (B.P. 165°C), (+)-form of which is found in oil of savin.

Sabinol, C₁₀H₁₆O. The 2-hydroxy derivative of sabinene found in the oil from *Juniperis phoenicea*. A colourless oil, B.P. 208°C (+)-form.

Saccharic Acid, C₆H₁₀O₈. The (+)-form crystallises in colourless needless, M.P. 125-126°C. Soluble in water, prepared by oxidizing starch or glucose by HNO₃.

Saccharin, C₇H₅NO₃S, $\begin{bmatrix} O \\ \parallel \\ S \\ O_2 \end{bmatrix}$. White crystals, M.P.

224°C (decompose). Obtained by oxidizing toluene-o-sulphonamide with alkaline permanganate. Used as a sweetening agent as its Na-salt (550 times more sweet than surcose).

Sachase Reaction. A method for the manufacture of ethyne from natural gas.

 $2CH_4 \xrightarrow{1500^{\circ}C} C_2H_2 + 3H_2$

Sacrificial Protection. A process of protection against electrolytic corrosion, e.g. Zn protects iron pipes buried under ground.

Safrol, shikimole, C₁₀H₁₀O₂. Colourless liquid, M.P. 11°C, B.P. 232°C. A chief component of oil of sassafras.

Sal Ammoniac. NH4Cl.

Salicin, C₁₃H₁₈O₇, C₆H₁₁O₅ O. C₆H₄CH₂OH. (β-D-glucoside of o-hydroxy benzyl alcohal). Colourless crystals, M.P. 201°C. Occurs in willow and poplar.

Salicyl Alcohol, Saligenin, 2-hydroxybenzyl Alcohol, C7H8O2 M.P.

87°C Prepared by reducing salicylaldehyde.

Salicylaldehyde, 2-hydroxybenzaldehyde, C₇H_eO₂. An oily liquid, B.P. 196°C. Obtained from CHCl₃, caustic potash and phenol (The Reimer-Tiemann reaction).

Salicylic Acid, 2-hydroxybenzoic Acid, C₇H₆O₃. M.P. 159°C. Occurs as oil of wintergreen (methyl salicylate). Manufactured by heating NaOC₆H₅ with CO₂ (under pressure) followed by acidification. Used in the manufacture of dyes, pharmaceuticals rust releasing fluids, etc.

Saline. A salt such as sodium chloride.

Salol, Phenyl Salicylate, C₁₃H₁₀O₂. M.P. 43°C. Obtained from NaOC₆H₅ plus sodium salicylate in the presence of POCl₅. Used as an enteric coating for pills, as a paint for throat ail ments, as an antiseptic, etc.

Salt A substance obtained by the interaction of equivalent quantities of acid and base, e.g., NaCl (normal salt), NaH SO₄ (acid salt), etc.

Salt Common. Sodium chloride.

Salt Bridge. A device (U-tube) used to reduce liquid-junction potential in an electronchemical cell containing a saturated solution of a salt (generally KCl) in an organic gel, e.g. agar. The

- Salt Cake (Sodium sulphate). An intermediate product in the manufacture of sodium carbonate by Leblanc process (common salt plus H₂SO₄).
- Salt Hydrates. Clathrates in which metal ions are surrounded by a fixed number of water molecules. The water molecules are bound to the metal ions by ion-dipole interations.
- Salting Out. A term used for the coagulation of hydrophilic sols. (colloidal solutions) and the reduction in solubility of a non-electrolyte by addition of a strong electrolyte (non-colloidal solutions), e.g., salting out of soaps and addition of NaCl to surcose to reduce its solubility in water.
- Salt Volatile. Ammonia, ammonium carbonate, oil of lemon and nutmeg in an alcoholic solution. Used as a first aid stimulant.
- Samarium Sm. At.No. 62, At. wt. 150.36, M.P. 1072°C, B.P. 1778°C, D. 7.52. A silvery metal of lanthanide series. Used in permanent magnets (SnCO₃), in optical glasses and in nuclear industries. Shows +3 and +2 oxidation state.
- Sand. A mineral resulting from rock disintegration.
- Sandmeyer's Reaction. A method for the synthesis of chloro and bromoarenes by heating the diazonium salts with Cu(I) halide
- Sandwich Compounds. A name applied to the bis-π-cyclopentadienylmetal compounds such as ferrocene. A type of complex formed between transition-metal ions and aromatic compounds in which the metal ion is 'sandwiched' between the rings. Bonding is between 'd' orbitals of metal and π electrons of ring.
- Santene. C₉H₁₄. Colourless oil, B-P. 140°C. A terpene like hydrocarbon (only) present in sandalwood oil and in pine-needle oil.
- Santenone, π Norcamphor. C₉H₁₄O. Solid, M.P. 56-59°C. Camphor like smell. Found in sandalwood oil. Prepared by the oxidation of santene with chromic acid (optically inactive form).
- Saponification. Hydrolysis of an ester with an alkali (NaOH).

RCOOR'+NaOH → RCOONa+R'OH Sod. salt

Saponins. Glycosides characterized by forming colloidal aqueous solutions which foam on shaking. Present in plants used as fish poisons.

- Sapphire. A blue form of ammonium oxide (corundum, a gemstone) having some metal impurities, prepared by fusing Al₂O₃. Used for handling HF.
- :Sarcosine, N Methylglycine. C₃H₇NO₂, CH₃NHCH₂COOH, M.P. 213°C (decomp.).
- Satin White. A mixture of CaSO₄ and Al(OH)₃ [Al₂(SO₄)₃ or alum plus Ca(OH)₂]. Used as a pigment.
- Saturated Compound. An organic compound in which atoms are linked by single bonds and does not contain any double or triple bonds, e.g, methane (CH₄), propane (C₈H₈), etc.
- Sawhorse Projections. (Fischer Projections). Describing the relative locations of the ligands (atoms, groups) attached to adjacement atoms.
- Sb. Antimony.
- s-Block Elements. Elements having the electronic configuration ns¹ or ns² [excluding elements in which (n-1)d levels are occupied]. e.g., I A (H, Li, Na, etc.), II A (Be, Mg, Ca, Sr, etc.) group elements.
- SBP Spirits. Special boiling point spirits.
- SBR. Styrene butadiene rubber.
- Sc. Scandium.
- Scandium, Sc. At. No. 21, At. wt. 44.96, M.P. 1539°C, B.P. 2832°C, D 3.0. A silvery element of group III with electronic configuration 4s² 3d². Its important ores are thortveitite (Sc₂Si₂O₇) and monazite. Shows only +3 oxidation state. Used in high
- Scavengers. Additives (e.g., tetraethyl lead or tetramethyl lead) which act as anti-knick to gasoline, lead scavengers must be used to ensure that lead oxide is not deposited in combustion.
- Schaffer Acid, C₁₀H₈O₄S (2-hydroxy-7-naphthalene sulphonic acid).

 A dyestuff intermediate. Prepared by sulphonation of 2-naphtol
 with H₂SO₄.
- Scheelite CaWO4. A tungsten mineral.
- Schiff's, Base. Compounds obtained by reacting aromatic amines with aliphatic or aromatic aldehydes and ketones. N-arylimides or R₂C=NAr compound, which form a secondary base on reduction with Na/C₂H₈OH. Weakly basic compounds.
- Schiff's Reagent. An aqueous solution of magneta (rosaniline) decolorized with H₂SO₃. A test for aldehydes and ketones. Aliphatic aldehydes and aldose sugar form a megneta colour with Schiff's regent (aromatic ketones do not react)

Schomaker-Stevenson Equation. An empirical equation

 $r_{A-B} = r_A + r_B - 0.09(X_A - X_B)$

where r_{A-B} is the bond length, r_A and r_B are the radil of A and B atoms respectively, and XA and XB the electronegativities of two atoms.

- Schonite, K2SO4, Mg SO4, 6H2O. A mineral of stassfurt salt. A potential source of K2SO4.
- Schotten-Baumann Reaction. A method for the preparation of N-phenylbenzamide (benzanilide) from a mixture of phenyl amine (aniline) and sodium hydroxide and benzyl chloride (stirring).

 $C_6H_5NH_2+C_6H_5COC1 \longrightarrow C_6H_5NH.COC_6H_5+HC1$

N-phenyl-benzamide is precipitated.

Schottky Detect. Defect structures.

Schradan. [(Me)₂N]₂ P(O) OP(O) (N Me₂)₃. A toxic to insects.

Schrodinger Wave Equation. An equation co-relating energy to field in wave mechanics.

Schweizer's Reagent. A dark blue solution formed by dissolving Cu(OH)₂ in concentrated ammonia solution. Used for the manufacture of rayon and as a solvent for cellulose.

Scintillation Counting. A process of detection and estimation of radioactivity by the scintillation produced when the ionizing radiation interacts with scintillation medium.

Screening. Separation of materials having particles of different sizes into fractions containing particles whose size lies within a certain range, by using sieves.

Scrubbers. Agents to remove impurity from a gas by washing it with a liquid.

Seaweed Colloids. The polysaccharides used in foodstuffs and textiles, e.g., agar algin, etc. (extraction of algae).

Sebacic Acid, Decanedioic Acid. HOOC. (CH2)8 COOH. M.P. 134°C. Obtained from castor oil and alkalis. Esters used as plasticizers.

Second. A base unit of time.

Second-Order Reaction. A reaction whose rate equation involves two concentration terms or the sum of the powers to which concentration terms are raised is two. E, g. Hydrolysis of esters by alkalis. The time required for the completion of a fraction of the reaction is dependent on initial concentration of reactant(s).

- Secondary Radiation. Radiation produced by the absorption of some other radiation (more strong), e.g., X-rays produced when X-rays strike a body.
- Sedimentation. The settling of a suspension, either under gravity or in a centrifuge. Used to estimate the average size of particles, in ultracentrifuge to find the relative molecular weights of macromolecules.
- Seed Crystal. A small crystal which when added to a supersaturated solution facilitates crystallisation of the whole solution.
- Segregation. A process by which solid particles (dry or suspension) arrange themselves in various layers according to size.
- Seignette Salt, C₄H₄O₆KNa, 4H₂O. Sodium potassium tartrate.
- Selection Rule. A rule controlling the transition between electronic energy levels in species.
- Selenates. Salts of selenium containing (SeO₄)²⁻, (HSeO₄)⁻ ions.
- Selenic Acid. H₂SeO₄. A strong acid (oxidation of selenates (IV) by Cl₂ or MnO₄), loses O₂ on heating.
- Selenides. Binary compounds of Se with other elements. Similar to sulphides and hydrolysed to H₂Se.
- Selenites. Salts of Se containing (SeO₃)²⁻ ions.
- Selenium Se. At. No. 34, At. wt. 78.96, M.P. 217°C. B.P. 684.9°C, D. 4.79. A metalloid element existing in several allotropic forms and belonging to VI group of periodic table. Electronic configuration 4s²4p⁴. Occurs in small amounts in sulphide chains with some cross linking. The common grey allotrope is sensitive metal).
- Selenocyanates. Derivatives of Se containing the Se CN grouping (eg, KSeCN from KCN+Se).
- Self-Ionization. A process of partial dissociation into cations and anious (self-ionizing solents)

 $2H_2O \Rightarrow (H_3O)^+ + OH^-$

- Semicarbazide, Aminourea. H₂N.CO.NH.NH₂.CH₅N₃O. M.P. 96°C. Forms semicarbazones with carbonyl compounds. Used for preparation and identification of aldehydes and ketones.
- Semi-Carbazones. A class of organic compounds containg the group by reacting an aldehyde of

- ketone in alcohol, dil CH3COOH or pyridine with semicarbazide (H₂N.NH CO.NH₂). The compounds have sharp melting points and are used as derivatives for the identification of aldehydes oriketones.
- Semi-Conductors. Materials containing impurities or imperfections resulting in the temperature dependent conductivity. This is due to the fact that the highest occupied energy level is very close to an unoccupied level, e.g., Zn O. Used in electronics, e.g, transistors, microprocessors, rectifiers and thermistors, etc.
- Semi-Permeable Membrane A membrane (natural or synthetic) which is permeable to some substances, e.g., water but does not allow others, e.g., salt, ions, etc. E.g. Parchment, (viz., bladder (natural) or copper cyanoferrate (synthetic), etc.
- Sephadex. A trade name for hydrophilic substance obtained from dextran. Used in get filtration.
- Sequestering Agent. (Chelating agents). Describing compounds which are very effective in forming complexes with metal cations, thereby, preventing them acting as simple hydrated cation, e.g., gluconic acid. Used as chelating agents.
- 2-Amino 3-Hydroxy Propionic Acid. C₈H₇NO₃, CH₂OH. CHNH₂.COOH. Colourless, M.P. 228°C (decomp.)
- Sesqui. (Oxide). Prefix indicating a 2/3 ratio, e.g., (M2O3) or Fe2O3.
- Sessile Dislocation. Describing a dislocation in a crystal lattice which can not glide.
- Sewage Treatment. A process by which waste water from industries or domestic source is filtered to remove solids and colloidal particles (using micro-organisms) and gases such as methane are used as fuels.
- Sextet. A series of six closely-spaced transitions in a spectrum.
- Shale Oil. An oil obtained by heating shales at about 950K.
- Shear Structures. Interrupted structure of crystals, e.g., heavy metal
- Shellac. A resinous secretion of the insect. Tachardic Lacca. Used as a resin.
- Describing a group of electrons that share the same principal quantum number, e.g., K-shell (n=1), L-shell (n=2) and M-shell (n=3).
- Sheradizing, Vapour Galvanizing. A technique used for coating metal articles (iron) by heating with Zn dust in a closed cessel below the M.P. of Zn.

Shift Reaction. Describing a reaction in which CO reacts with steam to form H₂ and CO₂

 $CO+H_2O$ Co catalyst CO_3+H_3

Process is used in oil or solid fuel gasification for the manufacture of 'towngas'.

Shift Reagents. Paramagnetic species which can shift appreciably nuclear magnetic resonance (n.m.r.) in any species with which they form weak complexes.

Si. Silicon.

Sialic Acid. N-and O-acyl derivatives of neuraminic acid.

Sialons. Si-Al-O-N systems. Used in machine tools.

Side-Reaction. A reaction that takes place simultaneously with the main chemical reaction.

Siemen's Process. A process for the manufacture of steel in which scrap metal, iron oxides and ferromanganese are added to molten pig (cast) iron.

Sigma Bond, ∂ -Bond. A covalent bond formed between the two electrons by the overlapping of the orbitals forming the bond being in one region in space directly between the nuclei.

Siemens (mho), S. The SI unit of electrical conductance equal to one ohm⁻¹.

Silanes. Hydrides of silicon, e.g., SiH₄, Si₂H₆, Si₃ H₈, etc. Obtained by the action of acids of Mg₂Si (magnesium silicide). Unstable compounds, ignite in air spontaneously.

Silatranes. Internal complexes of silicon

Silazanes, Compounds of Si having Si-N bonds.

Silica, SiO₂ (Silicon dioxide). An important constituent of earth's crust. Polymorphic, occurs as quartz (to 573°C), tridymite (to 1470°C) and cristobalite (to1710°C). Sand is mostly SiO₂. Naturally occurring silica is coloured due to Fe. Common flint is amorphous silica. Quartz crystals are non-centro-symmetric and can be right-or left-handed. Used in the manufacture of glass.

Silica Gel. An amorphous form of hydrated silica made by coagulating sodium silicate sol or by decomposing some silicates.

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The gel on standing, sets into a jelly. The dried gel is used as a catalyst support and as a drying agent. Used (dry gel) as an adsorbent in the recovery of solvents, for drying air, drying agent, for refining mineral oils, etc. Gel incorporated with cobalt cocompounds is used to remove moisture (pink-moist, blucdry).

- Silicates. Compounds containing metal ions and complex silicon oxygen compounds. The negative ion in silicates are of the type SiO₄⁴, Si₂O₇⁶, etc. Containing SiO₄ units linked in long type SiO₄⁴, Si₂O₇⁶, etc. Containing SiO₄ units linked in long type SiO₄⁴, Si₂O₇⁶, etc. Containing SiO₄ units linked in long type SiO₄⁴, Si₂O₇⁶, etc. Containing SiO₄ units linked in long type SiO₄⁴, Si₂O₇⁶, etc. Aluminosilicates chain sheets or three dimensional arrays. e.g., Aluminosilicates and borosilicates contain Al or B atoms in the structure. Natural silicates occur as rocks. Important silicates are feldspar, mica, zeolites, etc.
- Silica Vitreous, Quartz Glass. An amorphous vitreous material obtained on cooling fused silica. Used in optical works (low co-efficient of expansion).
- Silicic Acid. Unstable acid obtained by adding dilute sodium silicate solution to excess of dilute HCl followed by dialysis. Coagulates to a gel.
- Silicides. A compound of Si with a more electropositive element, e.g., Mg.Si.
- Silicon, Si. At. No. 14, At.wt. 28 085, M.P. 1410°C, B.P. 2355°C, B.D. 2.33. A hard brittle grey metalloid of group IV and the second abundant element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. Occurs as SiO₂ second element in the earth's crust. O
- Silicon Carbide, Carborundum, SiC. A black, very hard crystalline solid obtained by heating silicon (IV) oxide (sand) and C at 2000°C. Used for refractory bricks, crucibles, abrasive tools, etc.
- Silicon Rubbers, (Me₂SiO)_n. Derivatives of silicone having elastic properties over very wide temperature range. Inert to acids, alkalis, oils and oxidizing agents.
- Silicones. Polymeric synthetic silicon compounds containing chains of alternating silicon and oxygen atoms, with organic groups

bound to silicon atoms. Used as lubricating agents, water repellants and in waxes and varnishes. Have high flash points and are immiscible with water.

- Silicon Hydrides. See silanes.
- Silicon Nitride, Si₃N₄. Grey inert solid obtained by Si and N₂ at 1500°C. Resistant to acids.
- Silicon Oxide (mono), SiO. Formed by SiO₂ plus C in electric furnace. Used as a pigment and abrassive (Monex).
- Silk. A protein fibre obtained from the cocoons of the silk moth.
- Siloxanes. Compounds containing Si-O-Si groups with organic groups bound to the Si atoms, e.g., hexachlorosiloxane, Cl₃ SiOSiCl₃,
- Silver, Ag. At. No. 47, At. wt. 107.8680, M.P. 962°C, B.P. 2212°C, D. 10.49. An element (metal) of the copper group (I) with electronic configuration 4d¹0 5s¹. Occurs as the sulphide (Ag₂S), pyrargyrite or ruby silver, Ag₃ SbS₃. Extracted as a by product in refining copper and lead ores. It darkens in air due to the formation of silver sulphide. Shows oxidation states +4 to +1. Used in coinage alloys, table ware, and jewellery and its compounds are used in photography.
- Silver Bromide, AgBr. A pale yellow precipitate obtained by adding a soluble bromide solution to a solution of AgNO₃.

 $(AgNO_3+NaBr \longrightarrow AgBr+NaNO_3)$ M.P. 420°C.

- Silver Chloride, AgCl. A white solid, M.p. 449°C. Precipitated by the addition of a soluble chloride solution to a solution of AgNO₃. Occurs as 'horny silver'. Soluble in ammonia (aqueous) or sodium thiosulphate. Used in photography.
- Silver Halide Grains. The individual silver halide crystals in a photographic emulsion.
- Silver Nitrate, AgNO₃. Prepared by dissolving Ag in HNO₃ and crystallizing. M P. 212°C. On heating forms Ag, N₂O₄ and O₂.
- Silver Salt. The commercial name for sodium anthraquinone2-
- Silylation. A process for the formation of R₃Si-derivatives (R=H
- Silyl Compounds. Derivatives of the H₃Si-group.
- Simazine, 2 chloro—4,6—Bis (Ethylamino)—s-Triazine, C₇H₁₂ClN₅. A pre-emergency herbicide.

Simmons-Smith Reagent. A reagent discovered by du Pont chemists and having a formula

ICH.ZnI

- Single Bond. A covalent bond between two atoms involving one pair of electrons only. Often referred to as σ bond, represented by a single line, e.g., H-Br. Also, sometimes, applied to π bonds, e.g., Ni (P F₃)₄.
- Singlet. Describing a single transition in a spectrum with other levels at very different energies.
- Sinigrin, C9H16KO9S2. M.P. 126°C. A mustard glycoside.
- Sintering. A process of bonding by atomic or molecular diffusion in which powders are heated under pressure but at a temperature below the M.P. Sintered glass is a porous material used for
- Sizing. The process of filling the pores of paper and providing it some degree of water-repellency-either by the addition of rosins and alum (engine sizing) or by spraying the surface of paper
- Skatole, 3-Methylindole, C9H8N. M.P. 95°C. A volatile species present in coal tar or being formed in intestines by intestinal
- Slack Wax. An oily wax obtained from crude oil fractions by chilling or by the action of methyl-ethyl-ketone.
- Glass like compound or liquid solution of oxides formed during the extraction of metals (smelting and refining) when the impurities in an ore react with a flux. For example, in blast furnace the flux used in limestone (Ca CO₃) and the main Slag. impurity is silica (SiO₂). The slag formed is chiefly calcium $CaCO_3 + SiO_3 \longrightarrow CaSiO_3 + CO_3$ silicate:

Slaked Lime. Ca(OH):

Slurry. A thin paste of suspended solid particles in a liquid.

Slush Bath. Low temperature bath.

Sm. Samarium.

Smekal Defect. A crystal defect.

Smelting A process for extracting metals from their ores by melting it with other materials, such as C (for Zn and Sn) or CO (for iron). Cu and Pb are obtained by reduction of their oxide with the sulphide .

Smokeless Fuel. A fuel that produces minimum smoke when bu at in domestic fire place, e.g., a fuel obtained from low-temperature carbonization of coal.

Sn. Tin.

SNG. Substitute Natural Gas.

Soaps. Sodium and potassium salts of fatty acids (e.g., stearic, palmitic and oleic acids). Soaps are used to improve the cleansing properties of water.

Soda Ash, Na₂CO₃. Sodium carbonate.

Soda Lime. A grey material obtained from quicklime and NaOH solution followed by heating the product to dryness. Used as a drying agent and an adsorbent for CO2.

Sodamide. NaNH2.

Sodium Na. At. No. 11, At. wt. 22.987, M.P. 97.81°C, B.P. 882.9°C, D. 0.97. An alkali metal of I group with electronic configuration 2s1. Occurrs in sea water as NaCl. Prepared by electrolysis of fused NaCl or NaCl-CaCl2. A soft-silvery white metal, attacked by water (kept in kerosine oil). Used as a heat transfer medium, conductor, as a reducing agent.

Sodium Amide, NaNH2. Prepared from Na plus NH3. Used in

Sodium Ammonium Hydrogen Phosphate, Microcosmic NaNH4HPO4, 4H2O. Colourless, obtained from Na2HPO4 plus NH4Cl or (NH4)2HPO4 plus NaCl. Solidifies to glasses to produce specifice colours with some metal.

Sodium Antimonyl Tartrate. NaSbOC4H4O6. Antimony sodium

Sodium Azide. NaN3. A white solid obtained by passing NO over heated sodamide. Used as a reagent in organic synthesis and in lead azides (in detonators).

Sodium Benzoate, C6H5CO2Na. A white powder, soluble in water. Used as corrosion inhibitor and as an antiseptic.

Sodium Bicarbonate. See sodium hydrogen carbonase.

Sodium Bisumthate. NaBiO3. A very strong oxidizing agent prepared

Sodium Borate Na₂B₄O₇, 10H₂O

Sodium Bromide. NaBr. Prepared by Na2CO3 or NaOH plus HBr or Br₂ plus hot NaOH solution M.P. 757°C.

Sodium Carbonate. Na₂CO₃ (soda ash, Na₂CO₃). A white amorphous powder, manufactured by the ammonia soda or Solvay process

by Brine plus ammonia (NH₃) plus CO₂. NaHCO₈ is decomposed to Na₂CO₃ at 175°C. Forms decahydrates Na₂CO₅, 10H₃O when crystallized from aqueous solution (washing soda). Used in the manufacture of NaOH and washing soda. This salt is alkaline in solution due to hydrolysis:

 $CO_3^{2-}+H_3O^+ \rightleftharpoons H_2CO_3+OH^-$

- Sodium Chlorate, NaClO₃. A white solid formed by the action of Cl₂ on hot NaOH solution or by electrolysis of NaCl in water. A strong oxidizing agent. Used as an explosive, in pulp industry for bleaching (ClO₂) and as a herbicide.
- Sodium Chloride. NaCl. M.P. 801°C, D. 2.17. A white solid (common salt) obtained by HCl gas and NaCl solution (pure). Occurs in sea water (3%) Manufactured by solar evaporation of sea water. Solid NaCl is used in freezing mixtures, and preserve food and for preparing sodium carbonate (Solvay process). The sodium chloride crystal is cubic with octahedral co-ordination about Na and Cl.

Sodium Chromate (VI), Na₂CrO₄, 10H₂O.

Sodium Citrate, C₆H₅O₇Na₈, 2H₂O (also 5.5 H₂O). A citric salt. Used as a blood anticoagulant, to reduce blood acidity, and to prevent the formation of large curds in stomach of infants.

Sodium Cyanide, NaCN. M.P. 564°C. A poison.

Sodium Dichromate, Na₂Cr₂O₇, 2H₂O.

Sodium Dithionate, Na₂S₂O₄. A powerful reducing anent obtained by reducing Na₂SO₃ plus excess SO₂ with Zn. Used for removing O₂ from gases in presence of 2-anthraquinone sulphonate.

Sodium Ethoxide, Sodium Ethylate. A white solid obtained by neutralization of ethanoic acid with either sodium carbonate or NaOH. Used in organic synthesis and in dyeing industry.

Sodium Fluoride, NaF. A white solid (M.p. 902°C) formed by Na₂CO₃ or NaOH plus HF. Used as an antiseptic, an agent to prevent fermentation and as a constituent of ceramic enamels.

Sodium Glutamate, MSG (monosodium glutamate), C₈H₈N NaO₄, H₂O. A flavouring agent used as a food additive.

Sodium Hexafluoroaluminate, Na₃AlF₆. Cryolite.

Sodium Hydride, NaH. A white crystalline solid prepared by passing pure dry H₂ over Na at 350°C; Na is suspended in an inert medium. Reacts with water to form NaOH solution and H₂. Used as a powerful reducing agent to convert water to H₂, Conc. H₂SO₄ to H₂S and iron (III) oxide to iron. It bursts

- into flames spontaneously when comes in contact with halogens at room temperature. Dissolves in liquid ammonia to form NaNH₂ (sodamide).
- Sodium Hydrogen Carbonate, Sodium Bicarbonate, NaHCO₃. A white solid obtained by CO₂ and Na₂CO₃ solution. Sparingly soluble in water. Used in baking powder and in fire extinguishers.
- Sodium Hydroxide, NaOH. M.P. 318°C, D. 2.13. A white deliquescent translucent solid. Caustic Soda obtained by electrolysis of sodium chloride using a mercury cathode (Castner-Kellner cell). Dissolves in water with evolution of heat. Its solution is very corrosive and highly alkaline. Used in the manufacture of rayon (50%), paper (15%), petrochemicals, etc.
- Sodium Hypochlorice, NaOCl. Used as bleaching agent and an antiseptic (Cl₂ plus cold NaOH solution).
- Sodium Iodate, NaIO₃. I₂ and NaClO₃ form the salt. Used in volumetric analysis.
- Sodium Iodide, NaI. Formed from aqueous HI and Na₂CO₃ (M.P. 660°C).
- Sodium Lactate, C₃H₅O₃Na. Used in calico printing as a plasticizer for casein.
- Sodium Nitrate, NaNO₃, Chile Saltpeter. A white solid prepared from NaOH or Na₂CO₂ plus HNO₃. Used as a fertilizer and to prepare sodium nitrite.
- Sodium Nitrite, NaNO₂. A yellowish solid obtained by the thermal decomposition of sodium nitrate. Used for diazotization in organic reactions and as corrosion inhibitor.
- Sodium Perborate, NaBO₃, 4H₂O. Used in bleaching, cleaners, etc.
- Sodium Perchlorate, .NaClO₄. Prepared by heating NaClO₃. Used to form perchlorates.
- Sodium Periodates. Used as oxidizing agents. For example, NaIO₄, Na₅IO₆, etc.
- Sodium Peroxide, Na₂O₂. A yellowish white ionic solid obtained by burning Na in O₂. Used as a bleaching agent and an oxidizing meterial for wool and wood pulp.
- Sodium Phosphates. Several phosphates are known such as Na₂HPO₄ NaH₂PO₄ and Na₃PO₄, etc. Used as detergents, in food stuff and water conditioning.
- Sodium Potasium Tartrate, Rochelle Salt, Seignette Salt C₄H₄O₆ NaK, 4H₂O. Used as a saline aperient and reducing agent for silver mirrors.

- Sodium Pyrosulphite, Na₂S₂O₅. Used in photography.
- Sodium Saccharine, C7H4NNaO3, 2H1O. A sweetening agent.
- Sodium Silicates. Used in the manufacture of silica gel, as adhesive and in detergent manufacture. Various silicates are known.
- Sodium Stannate, Na2SnO3, 3H2O. A mordant (fuse SnO2 and NaOH).
- Sodium Sulphate, Glauber's Salt, Na₂SO₄ Commercially prepared from NaCl and H₂SO₄ at high temperature. Forms Na₂ SO₄ 10H₂O and Na₂SO₄, 7H₂O. Used in wood plup manufacture, glass and as a detergent.
- Sodium Sulphide, Na2S.
- Sodium Sulphide (poly). Na_2S_x (x=2,4, 5). Prepared from Na in liquid NH3 plus S.
- Sodium Tetraphenylborate, NaB (C6H5)4 Used for estimation of K.
- Sodium Thiocyanate, NaNCS. A volumetric reagent (NaCN+S).
- Sodium Thiosulphate, Na₂S₂O₃. A white solid manufactured by boiling sodium sulphite with flowers of sulphur or from the solid manufactured by boiling sodium sulphite with flowers of sulphur or from the solid manufactured by boiling solid ma Na₂SO₃ plus S. Used in photography (hypo) to dissolve silver halides. Also used as mordant and in volumetric analysis Crystallises from water as Na₂ S₂O₃, 5H₂O.
- Soft Detergents. Biodegradable detergents.
- Softeners. Materials added to rubber mix.
- Soft Soap. A liquid soap made by saponification with KOH.
- Sol. A colloidal consisting of solid particles distributed in liquid
- Solder. An alloy used in joint metals, e.g., a low melting barss. (M.P. 800°C) or Pb-Sn alloy.
- Solid Foams. A porous net work enclosing the gas (often in a liquidsolid system), e.g., charcoal.
- Solid Solution. A solution (solid) formed when two or more elements. or compounds share a common lattice, eg., some alloys
- Solid State Reactions. Reactions that take place between two or more solid compounds or decomposition occurring from the solid state, e.g., ternary oxides.
- Solubility The maximum amount of one phase (say solid) dissolved in the other phase (say liquid) at specified conditions of temperature and pressure. Solubility of gases is controlled by Henry's law.

Solubility Product. For a supersaturated solution of an electrolyte

 $AB_{solid} \rightleftharpoons AB_{dissolved} \rightleftharpoons A^{+} + B^{-}$ (solution)

The equilibrium constant for the reaction is

$$K = \frac{[A^+][B^-]}{[AB]_{solution}}$$

However the concentration of dissolved undissociated electrolute AB is taken as constant and so

 $[A^+][B^-]=K\varepsilon_p$

where K_{Ep} is the solubility produced of the electrolyte. The solubility produced exceeds if either an excess of A⁺ or B⁻ is added to the solution resulting in the precipitation of electrolyte.

Soluble Oil. Describing a mineral oil having emulsifiers which form stable emulsions when oil is added to water. Used in metal cutting and grinding process.

Solute. A substance added to a solvent to make a solution.

Solution. A liquid system of two or more species that are dispersed within each other at molecular level. Solutions are homogeneous and its components can be separated by altering the state of one of the components (freezing or boiling out one components). Its properties vary continuously with the proportions of components between certain limits. A dilute solution has small amount of solute. An ideal solution follows Raoult's low.

Solvation: The process of attraction of an ion (solute) by molecules of solvent. In general, a process of hydration.

Solvay's Process (ammonia soda process). An industrial process for making sodium carbonate. Also see sodium carbonate.

Solvent. In a solution, a liquid capable of dissolving other materials (solids, liquids or gases). The term is arbitrary except in cases such as chemical reaction between some constituents of a

Solvent Extraction. A process of dissolving out a substance from a material by shaking it with a suitable solvent.

Solvolysis. A reaction between a compound and the solvent in which it is dissolved, e.g., solvolysis of NH_4Cl to form free acid. $NH_4^+ + H_2O \rightleftharpoons NH_4OH + H^+$

Somatropie. A growth hormone.

- Sorbic Acid, 2, 4-Hexdienoic Acid, C₆H₈O₂, CH₃CH=CH-CH= CHCOOH. White solid, M.P. 134°C. Obtained by reacting (condensing) crotonaldehgde and ketone. Used as a food preservative.
- D-Sorbitol, D-Glucitol, C₆H₁₄O₆. M.P. 110°C. A alcohol corresponding to glucose. Prepared by reducing glucose in aqueous solution using H₂/Ni. Used for manufacturing ascorbic acid (vitamine C), in food stuffs, cosmetics, adhesives, etc.
- L-Sorbose, C₆H₁₂O₆. M.P. 160°C. Used as an intermediate in the preparation of ascorbic acid.
- Sour Products. Petroleum products such as gasoline or kerosine containing mercaptans.
- Soxhlet. An apparatus for the continuous extraction of a solid by a solvent.
- Space Lattice. An array of points in space in a crystalline structrure.

 These points may be occupied by atoms, ions or molecules.
- Spalling. The process of break-up of refractory with ultimate mechanical failure.
- Special Boiling Point Spirits (SBP). Refined soluent having specific selected boiling ranges and obtained by distillation from gasoline fractions, e.g., SBS 62/82°C. Used as solvents.
- Specic Conductance (K). The specific conductance is the conductance of the solution enclosed between two electrodes of 1 square centimeter area and are centimeter apart.
- Specifice Refractivity, r. Defined by

$$r = \frac{n^3 - 1}{n^2 + 2} \cdot \frac{1}{D}$$

where n=the retractive index of substance and D=the density of the substance.

- Spectral Sensitizers. The sensitizing dyes used to extend the natural sensitivity of a photographic silver halide emulsion. The process is known as spectral sensitization.
- Spectral Series. A group of related lines in the absorption of emission spectrum of a substance.
- Spectrochemical Series. Ligands arranged in order of their tendency to cause splitting at the d orbitals in complexes. The series is I-<Br-<Cl-<F-<OH-<H₂O <pyridine < NO₂.
- Spetrophotometer. An instrument for measuring different intensities of an electromagnetic radiation, in a spectrum, usually in the visible, infrared, or ultraviolet regin. It measures directly

(automatically) the relation between absorption of electromagnetic radiation and frequency (or λ).

Spectroscopy. The production and analysis of spectra. Used for analysis of mixtures, for identifying and determining the structure of chemical compounds, and for investigating energy levels in atoms, ions and molecules.

Sphalerite, (Zn, Fe) S. An ore of Zn.

Sphingomyelins. Phosphatides occurring in brain.

Spindle Oil. A lubricating oil of low viscosity.

Spin. A property af some elementary particles of spinning about an axis. In a magnetic field the spin lines up at an angle to the field direction and precess around this direction. The component of angular momentum along the field direction is $mh/2\pi$. For example, an electron has values $\pm \frac{1}{2}$.

Spinel. Mineral MgAl₂O₄.

Spinels. A class of mixed metal oxides represented by $M_2^{2+}M_2^{3+}O_4$ (where M=Mg, Fe, Co, etc., and $M_2^{3+}=A_1$, Fe, Cr, etc.)

Spin Moment. Defined as a magnetic moment produced when an electron spins on its own axis and is given by

 $\mu_0 = g \sqrt{S(S+1)}$

where μ_s =spin moment, g=gyromagnetic ratio and S=the sum of the spin quantum members of individual electrons.

Spin Quantum Number. Represents the clockwise and anticlockwise rotation of the electron. Two possible quantum numbers are $+\frac{1}{2}$ and $-\frac{1}{2}$.

Spin-Spin Coupling. A process of interaction between nuclear spins resulting in a fine structure in nuclear magnetic resonance spectra.

Spirans, Spiro-Compounds. Bicylic compounds containing only one atom common to both rings (spiro-atom).

Spirit of Salt. An obsolete name for hydro chloric acid.

Spodumene, LiAl (SiO3)3. A mineral of Li.

Spray Towers. A device in which the up going gas meets the down coming liquid sprays.

Squalene, C30H50O. An acrylic triterpene.

Sr. Strontium.

Stability Constant. An equilibrium constant for a process in which a complex in a formed in solution between a metal ion and a. ligand.

 $M+A \rightleftharpoons MA;$

 $K_1 = \frac{[MA]}{[M][A]}$

where K₁ is the stability constant and the terms in bracketts. represent activities. Also

- ∧G=RTln K₁

- Stabilizing Energy. The difference in energy between the delocalized structure and the conventional structure for a compound.
- Stabilizer. A substance added to prevent a chemical change.
- Stained Glass. A type of coloured glass prepared by adding impuri-
- Standard Electrode. An electrode which produces standard electromotive force, e.g., hydrogen electrode.
- Specifying standard conditions (e.g., 1 atmospherepressure, at 298 K temperature, etc.) for a chemical reaction Standard State.
- Standard Temperature and Pressure (STP). Specifying 1 standard atmosphere at 0°C. (1.013×10⁶ N/m² and 273.15K).
- Stannane. Tin Hydride, SnH₄. Colourless gas (SnCl₄+LiAlH₄ in etherate at -30°C) used as a reducing agent.
- Stannite, Cu2 Fe SnS4. A mineral.
- Starch (C₆H₁₀O₅)x. A polysaccharide that occurs exclusively in plants. Extracted from maize, barley, rice, potatoes, etc. Broken down by enzymes to simple sugars. Insoluble in cold water, but in hot water the granules gelatinize to form an opalescent dispersion. Used as an adhesive, for sizing paper and cloth, in volumetric analysis (indicator) and in foods, etc.
- Stassfurt Deposits. Describing deposits of various salts at Stassfurt
- Stationary Phase. A solid phase (e.g., alumina) having a large surface area which is used in chromatography.
- Stationary State. A state in a equilibrium reaction when the rates of forward and backward reactions are equal.
- Water vapours at temperatures above 100°C.
- Steam Distillation. A process of separating a liquid or a solid from other non-volatile impurities by distillat on in a current of other non-volatile impurities by distillation in a current of steam. Used for recovering, isolating or purifying substances.

The process is governed by the expression:

$$\frac{w_1}{w_2} = \frac{p_1 \mu_1}{p_2 \mu_2}$$

where w_1 and w_2 are the weights of the two components in the distillate and p_1 and p_2 are their respective partial pressures and μ_1 and μ_2 are their molecular weights respectively.

- Steam Reforming. The process of converting a mixture of methanesteam at 900°C into a mixture of CO and H₂ (in the presence of Ni catalyst).
- Steel. An alloy of iron and carbon (0.05-1.5%), eg, Alloy steel, Austenitic steel, Stainless steel (a group of Cr or Cr-Ni steels containing more than 12% Cr).
- Stereochemistry. Study of the spatial arrangement of atoms in
- Stereospecific Reactions. Describing reactions involving the breaking and formation of bonds at a single asymmetric carbon and resulting in a single stereoisomer.
- Steric Hindrance. Describing the influence exerted on a reacting group by the spatial arrangement of neighbouring atoms.
- Steroids. A general term applied to substances having the nuclear carbon skelton of the sterols, e.g., bile acids, sex hormones,
- Sterols. Alcohols having the following structure.

Stibine. Antimony hydride.

Stibinite, SbaS3. An antimony ore.

Stibite (Zeolite). 'Na₂Ca [(Al₂Si₆) O₁₆], 6H₂O. Used for softening

Still. A plant used for distillation,

Stoichiometry. The proportion in which elements form compounds.

A stoichiometric compound is one in which the atoms have com-

- Stopped Flow Spectrophotometry. A method to investigate the rates
- of fast chemical reactions. Streaming Potential. A potential difference caused by forcing a liquid with pressure through a diaphragm (reverse of electroosmosis flow).
- Strength of Acids and Bases. A measure of the ability of an acid to give hydroxonium ions (H₃O⁺) in aqueous solution or to accept protons by an aqueous base. E.g. H₂SO₄ (strong acid), KOH
- Streptomycia, C₂₁H₃₉N₇O₁₂. An antibiotic obtained from streptomyces medicinally griseus and used in its hydrochloride form. Used medicinally in the treatment of tuberculosis and in some infections due to organisms not susceptible to other antibiotics.
- Stripping. A process of separation of the more votatile component(s) in such of a liquid mixture from the less volatile component(s) in such a way that the less volatile component(s) is obtained in pure state but the more votatile component(s) may or may not be in pure state.
- Strontianite, SrCO₃. A mineral of Sr.
- Strontium, Sr. At. No. 38, At. wt. 87.62, M.P. 769°C, D 2.6. A soft metal of Group II with electronic configuration 5s². Occurs as strontianite (SrCO₃) and celestine (Sr SO₄) Manufactured by roasting the carbonate at 800°C followed by reduction with

$3SrO + 2Al \longrightarrow Al_2O_3 + 2Sr$

Also prepared by electrolysis of fused SrCl₂. Its compounds are used in pyrotechniques and flares, in glasses and in

- Strontium Carbonate, SrCO₃. A white insoluble solid (CO₃ over
- Strontium Chloride, SrCl₂. A white solid (Sr plus Cl₂ or Cl₂ over heated ScO). A specific plus to give red flame. heated SrO). Used in fire works to give red flame.
- A white insoluble salt occurring as mineral celestine. Prepared from SrO, Sr(OH)₂ plus H₂SO₄. Strontium Sulphate, SrSO4.
- Strychnine, C₁₂H₂₂N₂O₂. An important alkaloid of Nux vomica.

 Stimular N₂O₂. An important used to kill vermin. Stimulates all parts of nervous system, used to kill vermin.
- Styrene, Ethylene Benzene, Vinylbenzene, C₈H₈. Colourless aromatic liquid (B.P. 146°C). Polymerizes on heating to glassy resinous substanting. Manufactured substance which regenerates styrene on heating. Manufactured by all a leaved by all a leaved by all a leaved by all a leaved by dehydrogens. by alkylating benzene with ethene followed by dehydrogena-Used for manufacture of polymers such as polystyrene, ABS plastics etc.

- Styrene-Butadiene Rubber, (SBR). A synthetic rubber containing 25% styrene, 75% butadiene.
- Styrene Polymers. Polymers obtained by polymerization of styrene, e.g., polystyrene. Used for making films, as latex and as thermoplastics.
- Sub-boric Acid, (HO)2BB(OH)2.
- Suberane, C₇H₁₄. A trivial name for cyclopentane (B.P. 118°C).
- Sublimation. The conversion of a solid into a vapour phase without passing through the liquid phase, e.g., iodine.
- Sublimation Temperature. The temperature at which the vapour pressure above a solid is equal to the external pressure.
- Submicron. A term to define particles which are visible in ultramicroscope (10 m μ to 1μ).
- Substitution Reactions. Describing reactions involving replacement of one atom or group in a molecule by another atom or group, e.g., the formation of chlorobenzene (C₆H₆Cl) by the chlorination of benzene (C_H) nation of benzene (C₆H₆). Important substitution reactions are: Nucleophile Substitution (S_N): The attacking substituent is a nucleophile.

Electrophile Substitution: The attacking substituent is an

Free Radical Substitution: A free radical is the attacking

Substrate. A term used in enzymic reactions, the molecule on which

Succinic Acid, Butanedioic Acid, C₄H₆O₄, {CH₂-COOH} 182°C. Occurs in plants such as amber, sugar cane, algae, etc. Manufactured by the catalytic reduction of maleic acid. Forms anhydride, used in the manufacture of polyesters.

Succinic Anhydride, C4H4O3, CH2CO CH₂CO . White crystals, M.P.

120°C. Manufactured by heating succinic acid at 235°C. Used in the manufacture of dyest. in the manufacture of dyestuffs and polymers.

Succinamide, C₄H₅NO₂, (CH₂)₂C(O) NH.C(O), M P. 126°C, obtained by heating ammonium succin by heating ammonium succinate. Used for disinfecting water. Sucrase. An enzyme.

Sucrose (Cane sugar), C₁₂H₂₂O₁₁. Extracted from sugar cane and sugar beet. A disaccharide formula from sugar cane and and a sugar beet. A disaccharide formed by a glucose unit and a fructose unit. No reducing action, hydrolysed to glucose and fuctose by acids.

e by acids.

$$C_{12}H_{22}O_{11} + H_2O \xrightarrow{\text{acid}} C_6H_{12}O_6 + C_6H_{12}O_6 \text{ fructose}$$

It has 8 hydroxyl groups.

Sugar of Lead. Pb (O2C CH3)2, 3H2O.

Sugars. A term applied to carbohydrates which are crystalline, very soluble in water and generally sweet.

Sulphacetamide, C₈H₁₀N₂O₃S. M.P. 181-183°C. Used medicinally for treating urinary infections.

Sulphadiazine 2-(p-Aminobenzenesulphonamido)-pyrimidine,

 $C_{10}H_{10}N_4O_2S$. White powder, M.P. 255-256°C.

Effective against streptococci, staphylococci and other

Sulphaguanidine, p-Aminobenzenesulphonylguanidine, White powder, M.P. 189-190°C. Used for treatment of intes-

Sulpha merazine C₁₁H₁₂N₄O₂S. M.P. 236°C. Used in blood treat-

Sulphamethoxazole, C₁₀H₁₁N₃O₃S. Colourless M.P. 197°C.

Sulphamic Acid, H₃N⁺SO₃⁻. A strong acid soluble in water (SO₂) plus acetoxime). Used as an acidimetric primary standard,

Sulphanes. A group of hydrogen sulphides H₂S₂ to H₂S₆.

Sulphanilic Acid, 4-aminobenzene sulphonic acid. Prepared by eating aniline sulphate for 8-10 hours at 190°C.

Sulphates, SO₄². Derivatives of sulphuric acid.

Sulphathiazole, C₉H₉N₂O₂S₂. Active against β-haemolytic strepto-

Sulphide (S²-). A compound of S with other electropositive element,

e.g., Na₂S. Polysulphides contain (S_x²-) ion.

Sulphinyl. Compounds containing the group > SO Sulphite. A salt of sulphurous acid containing (SO₃)² - species.

Salpholane, C₄H₈O₂S, (|S|). A viscous liquid (B.P. 285°C)

Sulphonamides. Compounds having the grouping $-SO_2N <$. which solidifies to a colourless solid.

Sulphonation. A process of introducing sulphonic groups (-SO₂OH) into organic compounds.

Sulphones. Organic compounds with >SO2 grouping.

Sulphonic Acids. Organic compounds containing the group -SO₅OH, e.g., benzene sulphonic acid, C6H5SO2OH.

Sulphur, S. At No 16, At Wt. 32.06, M.P. 112.8°C, B P. 444.4°C, D 2.07 A POP 14 Pop 16 of Pop 16 D 2.07. A non-metallic solid, yellow coloured and member of Group VI with classic solid, yellow coloured and member as Group VI with electronic configuration 3s², 3p⁴. Occurs as sulphide ores (FeS.) sulphide ores (FeS₂) and sulphate rocks (CaSO₄). Exists in various attotronic modes. various attotropic modifications such as α-S, rhomatic sulphur, monoclinic sulphur, a lement monoclinic sulphur, β -sulphur, S_{λ} , S_{μ} , etc. A reactive element which combines S_{λ} , S_{μ} , etc. A reactive element which combines with most other elements. Shows oxidation states +6, +4 +2 0 most other elements. states +6, +4, +2, 0 and -2. Behaves as a non-metal with a great tendency to catenation.

Sulphur Dyes. Dyestuffs applied from solutions containing S⁻², SH
or polysulphide and applied from solutions containing S⁻², SH or polysulphide and are applied in reduced state and may or

Sulphuretted Hydrogen, H2S. Hydrogen sulphide.

Sulphuric Acid, H₂SO₄, S(O)₂(OH)₂. M.P. 10°C, B.P. 240°C, D 1.83. Manufactured from SO. Manufactured from SO₂ plus air over a Pt or V₂O₅ (500-600°C) then 400-450°C then 400-450°C. A solution of SO₂ in H₂SO₄ is fuming sulphuric acid. A powerful dehydrating agent. Handled in ((NH₄)₂SO₄), explosives pigments. ((NH₄)₂SO₄), explosives, pigments, petrochemicals, etc.

Sulphuric Acid, Fuming. A solution of SO₃ in H₂SO₄ containing

Sulphur-nitrogen Derivatives. Some derivatives such as S₄N₄ or polymeric derivatives (SN)x, S₇NH, S₆(NH)₂ etc.

Sulphurous Acid, H₂SO₃, SO(OH)₂. SO₂ in water forms SO₂. 7H₂O₇ a gas hydrate, and free H2SO3 does not exist.

Disulphur Monoxide, S2O. An unstable oxide.

Sulphurdioxide, SO₂. A colourless gas with typical odour. Formed by burning S metal sulphida gas with typical odour. by burning S, metal sulphides, H₂S in air or by the action of an acid on H₂S or sulphite. A powerful reducing agent. Solution behaves as an acid. Used for manufacturing of SO₃.

Sulphur trioxide, SO₃. A gas readily absorbed by water to form H₂SO₄ or by sulphuric acid to produce fuming sulphuric acid

(SO₂ plus O₂ - contact process).

Sulphur Oxyacids. Important oxy acids of S are as follows:

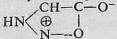
Thisulphuric acid Dithionous acid Polythionic acid Pyrosulphuric acid Peroxymonosulphuric acid

S(O) S(OH)2 O(HO) SS (OH)O O(HO) SSnS (OH)O O2(HO) SOS (OH)O2 O2S(OH) (OOH)

- Sulphuryl. Compounds of >SO2 group.
- Sulphydral (thio). The -SH group.
- Super Acids. Mixtures of mainly fluorine compounds, e.g., HF-SbF₅-Strong acids.
- Supercooling. A process of cooling a liquid to a temperature below that crystallization would occur without the separation of solid.
- Super Heavy Elements. Post actinide elements.
- Superoxide Dismutase (SOD). An enzyme which brings about the following reaction:

 $2O_2^- + 2H^+ \longrightarrow H_2O_2 + O_2$

- Superoxides. Compounds having the paramagnetic ion, O2-.
- Superphosphates. A mixture of calcium hydrogen phosphate, Ca(H₂PO₄)₂ and calcium sulphate. Used as a fertilizer.
- Surface Active Agents. Substances (soluble) which reduce the surface tension of liquids even in small concentrations (mainly organic)...
- Surface Combustion A process of burning gaseous fuel and air in a way so that catalytic oxidation occurs at the surface of a porus refractory.
- Surface Energy. The work done to increase the surface by unit area against the force of surface tension (in a system).
- Surface Tension. The force in dynes per unit length acting at right angles to the line along the surface of the liquid.
- Surfactant. A substance that lower surface tension, e.g., long chain sulphonic acids.
- Surgical Spirit. Methytated and denatured spirit by the addition of caster oil (2.5%), diethyl phthatate (2.0%) and methyl salicylate (0.5%).
- Suspension Agents. Materials used to hold up the solid in a suspension, e.g., Bentonite.
- Sweating. A process of removing oil and waxes (low melting) from the cake in dewaxing processes.
- Sweetening. A process of removing objectionable odour materials from gasoline and kerosine fractions of petroleum.
- Sweetening Agents. Substances used for sweetening purposes, e.g., sugars, saccharine, etc.
- Sydnones. Heterocyclic compounds represented as



Sylvestrene, C₁₀H₁₆. A monocyclic terpene.

Sylvine Sylvity, KCl. A source of K.

Sylvinite, KCI-NaCl. A source of K-salt.

Symmetry Elements. A term applied to crystals.

Axis of Symmetry. An imaginary line drawn through the centre of the crystal such that on rotation through 360° about this line, the crystal appears unchanged twice (2-fold), thrice (3-fold), or 4 times (4-fold).

Plane of Symmetry. An imaginary plane passing through the centre of a crystal and bisects the crystal in such a way that one part is the mirror image of the other.

Centre of Symmetry. A point through which there is reflection to an identical point in the pattern.

Syneresis. A process for separating liquid from a gel on standing.

Synergist. A process in which a substance (itself less active) enhances the activity of another substance present in the system.

Synthesis. Formation of complex compounds from simpler ones.

Synthesis Gas. A mixture of CO and H₂ produced by steam reforming of natural gas

$$CH_4 + H_2O \longrightarrow CO + 3H_2$$

T

- T. Tritium.
- 2,4,5-T, 2,4,5-Trichlorophenoxy Acetic Acid, C₈H₅Cl₈O₈. M.P. 155°C.
- Ta. Tantalum.
- Tactosols. A type of colloidal sols containing non-spherical particles and which are capable of orientating themselves.
- Talc. Mg₃ (OH)₂ Si₄O₁₀. A lubricant (French chalk). A silicate containing layers of linked SiO₄ tetrahedra (electrically neutral layers). Used as a filler in paints, rubber, insecticides, a constituent of toilet powder, etc.

Tannase. A class of moulds containing an enzyme which splits tannins (ester type), e.g., Aspergillus.

Tannic Acid, Gallotannic Acid, Tannin, A yellowish powder (M.P. 210-215°C).

R=Galloyl m-digalloyl, m-trigalloyl

Tanning. A process of changing skins and hides into useful leather.

Tanning. A class of amorphous substances present in plants, obtained by extracting the raw materials with water or other solvents and precipitating with lead ethanoate. Gives a blue-green colour with iron salts. Used in the treatment of hides to make leather and used as a mordant in the textile industry.

Tantalum, Ta. At. No. 73, At.wt. 180.948, M.P. 2996°C. D. 16.6. A transition element of Group V with electronic configuration 6s2 5d3. Prepared by reduction of Ta2 O5 with alkali metals or by electrolysis of fused fluorides. Used in surgical appliances.

Tapiolite. Fe Ta2O6. A mineral of Ta.

Tar. A non-aqueous liquid condensate obtained from the destructive distillation of carbonaceous materials.

Tar Acids. Compounds mainly of phenols. Obtained from tar (tar+Na₂CO₃ followed by CO₂ reaction).

Tar. Oils. Phytotoxic oils containing aromatic hydrocarbons. Used to kill insect (eggs) in trees.

Tartar-emetic. Antimony potassium tartrate.

Tartaric Acid, HOOC (CH.OH)₂.COOH. A crystalline hydroxy carboxylic acid (2,3-dihydroxybutanedioic acid). M.P. 170°C. Occurs in two optically active and two inactive forms. Occurs in grapes. Manufactured from tartar argol and wine lees by precipitation as Ca-salt followed by action with H2SO4. Used in drinks, textiles, etc.

Racemic tartaric acid, (±) has M.P. 273°C and mesotartataric acid (1H₂O), M.P. 140°C (anhydrous).

Tartronic Acid, Hydroxy Malonic Acid, HO₃C.CH(OH).COOH. Obtained by heating dinitrotartaric acid in aqueous solution (M.P. 160°C).

314 Tautomerism-

Tautomerism. A phenomenon resustting from the migration of a hydrogen atom within a molecule.

Tau Value, τ. A measure of shift in nuclear magnetic resonance.

Tb. Terbium.

Tc. Technetium.

TCNE. Tetracyanoethylene.

Te. Tellurium.

Technetium, Tc. At.No. 43, At. wt. 98.906, M.P. 2172°C, D. 11.5. A transition metal produced by bombarding Mo, with neutrons and also by fission of U. All isotopes are radioactive. An element of Group VII with electronic configuration 4d⁶ 5s¹.

Teflon. Fluorine containing polymers.

TEL. Lead tetraethyl.

Telluric Acid, Te (OH)6 Obtained from Te in aqua regia and a

Teliurides. Binary derivatives of Te with other elements.

Tellurium, Te. At. No. 52. At. wt. 127°60, M.P. 449.50°C, D. 6.24. A silveny element of VI group with electronic configuration 5s² 5p⁴. Shows oxidation states -2, +2, +4 and +6. Used in minerals, in stainless steel (Poisonous compounds).

Telomerization. A process of polymerization or condensation of a few molecules of olefine to form low molecular weight polymers, e g., (C₂F₄)₄.

Temperature Scale. A practical scale for measuring temperature

Tempering. A process of reheating a quenched steel (200-553°C) to reduce hardness, tensile strength and relive internal strength.

Tetratogen. A chemical used to produce malfunction in the form of

Terbium, Tb. At. No. 65, At. wt. 158.93, M.P. 1360°C, D. 8.28. A soft malleable metal of lanthanide series. Shows +3 and +4 oxidation states. Used as a laser material (sodium terbium

Terebene. A mixture of terpene hydrocarbon obtained by the action

Terephthalic Acid, C₈H₆O₄ (1, 4-benzene dicarboxylic acid), C₈H₆O₄. A sublimative solid (M.P. 300°C) Used for producing

Ternay Compound. Describing a compound having three elements,

Terpenes. Strictly terpenes are volatile, aromatic hydrocarbons having the empirical formula C10H16. Broadly speaking, the term includes sesquitrepenes, C₁₅H₂₄, diterpenes, C₂₀H₃₂ and even high polymers. Terpenes are characteristic products of many varieties of vegetable life and important constituents of most or dants.

Terphenyl, (p-Diphenyl benzene). Used in heat transfer fluids.

1, 8-Terpin, Terpine, C10H20O2. A monocyclic dialcohol having cis and trans isomers.

Terpinenes, C₁₀H₁₆. Monocyclic terpene having α, β and γ isomers.

Terpineols, C10H18O. A monocyclic terpene alcohol obtained by dehydrating terpin.

Terpinolene, C10H16. A monocyclic terpene obtained by the action of acids on pinene.

Tetraborane, (10), BaH10. A boron hydride.

Tetracene, 1-(5-tetrazolyl)-4 Guanyl Tetrazene, Hydrate, C2H8N10O. A pale yellow solid used as detonator.

Sym.-Tetrachloroethane, Acetylene Tetrachloride, HCCl2 CHCl2. A colourless toxic liquid (B P. 146°C).

Tetrachloroethane, Perchloroethylene, CCl2=CCl2. B.P. 121°C. Used as a dry-cleaning agent (solvent).

Tetracyanoethylene, TCNE, (NC), C=C (CN)2. A sublimable solid

(M.P. 200°C). A strongest π-acid.

7,7,8,8—Tetracyanoquinonedimethane, TCNQ. Accepts an electron from suitable donors, forming a radical anion. Used for colorometric measurement of free radical percursors, construction of heatsensitive resistors.

Tetracyclines. A class of antibiotics obtained from streptomyces. Tetracyline has the structure:

They are broad spectrum antibiotics active against Grampositive and Gram-negative bacteria.

Tetraethyl Lead. TEL.

- Tetraethylpyrophosphate, TEPP, C₈H₂₀O₇P₂. An insecticidal used as a direct acaricide or aphicide.
- Tetrahedral Compound. A compound having four bonds directed towards the corners to a regular tetrahedron (bond angle about 109°) e.g., CH4.
- Tetrahydrocannabinol, C21H29O2. An active component of hashish (marihuna) found in Cannbis Sativa.
- Tetrahydrofuran, THF, C4H8O. A solvent for resins and used in tetrahydrofuran polymers.
- Tetrahydrothiophen, THT. Colourless mobile liquid (B.P. 119°C). Used to provide an odour to domestic gas supplies.
- Tetralin, 1,2,3,4 Tetrahydronaphthalene, C₁₀H₁₂. A colourless liquid used as a solvent for fats, oils and resins.
- Tetramethylsilane, TMS, (CH₃)₄ Si. Used as an internal reference standard for proton magnetic resonance spectroscopy.
- Tetrose. A carbohydrate containing 4 carbon atoms, e.g., 4-aldo-
- Tetryl, N-2, 4,6-Trinitrophenyl-N-methyl nitramine, C7H5N5O8. A pale yellow powder (M.P. 129°C). A stable explosive.
- Th.
- Thallium, Th. At. No. 81, At. wt. 204.383, M.P. 303.5°O, D. 11.85.

 A soft malleable greyish metal of Group II. Occurring in crookesite (Ag, Cu, Tl)-Se. Electronic configuration 6s² 6p¹. Its compounds are used in poisons, glasses and in electronic
- Theobromine, 3,7-Dimethyl Xanthine, C7H8N4O2. An alkaloid
- Thermal Analysis. Analytical process used for identification and investigation of the products of the action of heat on a compound or complex, e.g., DTA (differential thermal analysis) and DSC (differential scanning calorimetry).
- Thermal Cracking. A process to increase the gasoline yield from a
- Thermitit Process. A process of reduction by Al powder and a metal oxide (Fe₂O₃) used to produce heat for welding.
- Thermobalance. A balance in which a sample can be heated to
- Thermochemistry. A branch of chemistry involving the heat changes accompanying heat changes.

Thermodynamics. The study of heat and other form of energy and the various related changes in physical quantities.

Themodynamics, First Law of. The total energy of an isolated system remains constant. The energy is transferred from one form to another. Mathematically,

$$dq = dE + \omega$$

where dq=the heat transferred to the system, dE=the increase in internal energy of system and ω is the work done (external).

Thermodynamics, Second Low of. 'Heat can not of itself transfer from a colder to warmer body' or it is impossible for a self-acting machine, unaided by an external agency, to convey heat from a body at a low to one at a high temperature.

Thermodynamics, Third Law of. 'For a perfect crystal at absolute zero on the kelvin scale the entropy is zero'.

Thermogram. A plot of weight against temperature.

Thermoplastic Resins. Describing resins which get softened by heating and harden again on cooling.

Thermosetting Resins. Resins which on moulding become infusible and insoluble.

THF. Tetrahydrofuran.

Thiamine, Aneurine, Vitamin B₁. An anti-neuritic factor the absence of which causes beri-beri disease in human body. Destroyed by heating above 100°C. Important source is yeast, eggs and the germ of cereals.

Thickening. A method to recover a solid(s) in the concentrated form from a slurry or suspension.

Thin Layer Carmatography. See Chromatography.

Thio. Containing S.

Thiocarbonates. Salts of thicarbonic acids.

Thiocarbonic Acid, H2CS3. Used for destroying fungus phylloxera.

Thiochrome, C₁₂H₁₄N₄OS. A yellow solid (M.P. 227-228°C) extracted from yeast and prepared by oxidizing thiamine.

Thiocyanates. Salts of thiocyanic acid, eg., AgSCN. Used as photographic sulphur sensitizers.

Thiocyanic Acid, HSCN. Gas at room temperature but polymerizes. Thiocyanogen. (SCN)

Thiokols. Polysulphide polymers.

Thiols. A name for mercaptans.

Thiomolybdates. Bidentate ligands, e.g., [MOS₄]²-.

Thionic Acids, H2SnO6.

Thionyl. Compounds containing >S=O groups.

Thionyl Chloride. SOCl₂ Obtained by SO₂ plus PCl₅. Used for dehydration of metal chloride hydrates and for converting COH to CCl grouping. (A liquid).

A colourless liquid with smell that of benzene (B.P. 84° C). Prepared by heating sodium succinate with P_2S_5 . It contains mobile π electrons.

Thiosulphates. Salts containing (S2O3)2- ions.

Thiouracil, C₄H₄N₂OS. White powder used to decrease the activity of thyroid gland.

Thiourea, CH₄N₂S, S=C (NH₂)₂. A white crystalline solid (M.P. 172°C). Manufactured from H₂S plus cyanamide. A photographic sensitizer.

Thixotropy. A process of transformation of isothermal gel-sol by shaking or other means (mechanical only).

Thorium, Th. At. No. 90, At.wt. 232.04, M.P. 1750°C, D. 11.7. A toxic radioactive element of actinoid series. Occurs as thorite and thorogummite and monazite. Prepared by reducing Th F₄ by Ca. Attacked by hot water and dilute acids. Shows +4 state. Forms ²³³U (from ²³²Th) and Th O₂ (a catalyst in Fischer-Tropsch process). Used as an oxygen remover in the electronic industry.

Thorium Oxide. Th O2. A white salt.

Thortveitite, Sc₂Si₂O₇. A source of Sc.

- Threose, C₄H₈O₄. A tetrose sugar.
- Thrombin, An enzyme which catalyses the reaction of fibrinogen to fibrin in clotting blood.
- Thujane, C₁₀H₁₈. A dicyclic hydrocarbon, a colourless oil (mobile) B.P. 157°C
- Thujone, C₁₀H_{1n}O. A colourless oil having smell of menthol. Occurs as α and β thujones. A dicyclic ketone present in thuja, worm wood, etc.
- Thujyl Alcohol, C10H18O. A secondary alcohol present in worm wood oil.
- Thulium, Tm. At. No. 69, At. wt. 168.93, M.P. 1545°C, D. 9.332. A soft, ductile silvery element of lanthanoid series. Forms a series of +3 compounds (Tm³⁺-pale green). Used as a portable source of X-radiation after neutron irrdiation.

Thymine, 5-Methyluracil, 5-Methyl 2, 6-Dioxytetrabydropyrimidine, H C.H.N.O. N

A constituent of deoxyribose nuclei acid (M.P. 321-325°).

- Thymol, 2-Hydroxy-1-Isopropyl-4-Methyl-benzene, C10H14O. A constituent of oil of thyme, manufactured from piperitone. Used as mouthwashes and a limited antiseptic.
- Thymol Blue, Thymolsulphonphthaleien. A member of sulphonphthalein group of indicators. Works in two pH ranges, e.g., 1.2 (red) to 2.8 (yellow) and 8.0 (yellow) to 9.3 (colourless).
- Thymolphthalein. C₃₈H₃₀O₄. An indicator of pH range of 9.0 (colourless) to 10.5 (blue).
- Thyroid Hormone, Thyrotropine. A glycoprotein hormone which stimulates the thyroid gland (M. wt. abaut 10,000) to produce
- Thyroxine, C₁₅H₁₁I₄NO₄. A hormone of thyroid. Deficiency of hormone leads to goitre, myxoedema and cretinism while excessive secretion causes Grave's disease.
- Ti. Titanium.
- Tin. Sn. At. No. 50, At. wt. 118.70, M.P. 232°C, D. 7.28. A H_3C-C-H

H₃C-C-COOH

white lustrous metal of low-melting point of Group IV with electronic configuration $5s^25p^2$. Occurs as cassiterite or tinstone, SnO_2 , which is reduced with C. Metal has three allotropic forms, e.g., grey tin (a) stable below 13.5° C, a brittle powder, γ -Sn (stable above 161° C) and Sn (white) stable above 13.5° C. The metal is attacked by halogens and conc. acids. Shows oxidation state of +4 (SnCl₄) and +2 (SnCl₂). Used in the alloys formation in coating steel, in glass coatings and as fungicide.

- Tin Alloys. Forms various alloys such as Brittannia metal, fusible metal, Babbitt metal, bell metal, antifriction metal, bronze, gun metal and pewter.
- Tin (II) Chloride, SnCl₂ (Stannous chloride). A white solid (M.P.) 247°C) obtained from Sn and gaseous HCl. Acts as a reducing agent, combines with NH₃ and forms hydrates. Used as a mordant.
- Tin (IV) Chloride, SnCl₄ (Stannic chloride). A colourless fuming liquid, soluble in organic solvents (B.P. 114°C). Prepared from Sn and Cl₂. Dissolves S, P, Br₂ and I₂. Dissolves in conc. HCl to form (SnCl₆)²⁻ anion.
- Tincture of Iodine. Alcoholic solution of iodine containing iodine (2.5%) and KI (2.5%)
- Tin Hydrides, SnH₄ (-52°C), Sn₂H₆. Compounds of Sn having Sn-H bonds. Good reducing agent.
- Tinning. A process of coating of iron with tin. Used for manufacturing tin-cans.
- Tin (II) Oxide, ShO. A dark green or black solid obtained by precipitating the hydrated oxide from a solution of Sn (II) ions and dehydrating the product at 100°C. Forms complexes.
- Tin (IV) Oxide, SnO₂. Occurs in nature as cassiterite or tinstone. A colourless crystalline solid obtained as hydrate from Sn (IV) solution and dehydrated to SnO₂. Exists as hexagonal or rhombic crystals. Insoluble in water. Used as a 'putty powder' for polishing glass and metal.

Tin Salt, SuCl2, 2H2O.

Tin Stone, Sn'Q2.

- Tin (II) Sulphide, SnS. A grey solid obtained from Sn and S at 900°C.
- Tin (IV) Sulphide, SnS₂. A yellowish solid prepared from a solution of Sn (IV) by H₂S or Sn plus S. Forms a yellow solid (mosaic gold) when heated with NH₄Cl and S.
- Titanates. Compounds of Ti such as Ti (OR)4. Used as catalysts.

Tetanium Ti. At. No. 20, At. wt. 49, M P. 1660°C, D 4.5. Astransition metal occurring as ilmenite (FeTiO₃) and rutile (TiO₂). Obtained by reducing TiCl, with Mg. Group IV element with electronic configuration 342 4s2. Stable oxidation state is +4 (TiO₂). Ti is reactive at high temperatures. Used as a light weight construction material and its compounds such as TiO4 as, a white pigment.

- Titamium Alloys. Forms alloys with Fe Mn, Cr and Fe. Used in aircraft industry and for making engines.
- Titanium Carbide, TiC. A steel grey solid. Used an a deoxidizer in. steel manufacture and in tool tips.
- Titanium Dioxide, TiO, Occurs naturally as three crystalline forms:
 rutile (tetragonal), brookite (orthorhombic), and anatase
 rutile (tetragonal) (tetragonal). An important white pigment obtained by hydrolysis of purified TiOSO4 or TiCl4 or TiCl4 plus O2 through
- Titration. A process in volumetric analysis involving the addition of a solution of known concentration (titrant) to a solution of unknown concentration from a burette until the equivalent point or end point of the titrant is reached. Volumetric titration may or may not require an indicator. Electrical methods are also used for titrations.

Tl. Thallium.

TLV. Threshold limit value.

Tm. Thulium.

TNT. Trinitrotoluene.

Toad Venom. A poisonous substance secreated by the skin glands of

Tocopherols. The methylated derivatives of tocol, e.g., a-tocopherol $(C_{29}H_{50}O_2)$, β -tocopherol $(C_{28}H_{48}O_2)$, γ -tocopherol $(C_{24}H_{46}O_2)$ and y-tocopherol (C27H46Os).

Tolan, C6H5C = C.C6H5. Diphenyl acetylene.

Tollen's Reagent. An ammoniacal solution of silver oxide which is used as a test for aldehydes (formation of Ag-mirror).

Toluene, C₇H₈, (methyl benzene). A colourless (B.P. 111°C),

steam volatile and refractive liquid. Prepared from petroleum fractions (rich in naphthenes) by catalytic reforming in the

presence of H₂ Or by the carbonization of coal. Forms benzoic acid (by chromic acid or HNO₃), benzaldehyde (mild oxidation), TNT, trinitrotoluene (by vigorous nitration) and sulphonic acid (by sulphonetion). Used as a solvent, a constituent of high octane aviation and motor gasoline.

Toluene, 2, 4-Diisocyanate, TDI, C9H6N2O2, C6H4(NCO)2. A liquid (B.P. 120°C) prepared by nitrating toluene. Used to form polyurethane.

Toluidine, C7H9N.

o-Toluidine, 2-aminotoluidine. Colourless liquid (B.P. 198°C). Obtained by reducing 2-nitrotoluene. p-Toluidine, 4-aminotoluidine. Colourless leaflets (M.P. 45°C). Prepared by reducing 4-nitrotoluene by Fe/HCl.

Toluyl. A group CH3C6H4CO-.

Tolyl. The group CH3C6H4CH2-.

Topaz, Al₂SiO₄ (F, OH)₃. Used as gemstones (transparent crystals)

Tor. A pressure unit and 1 torr is equal to 1 mm of Hg pressure.

Tosyl. Esters of toluene sulphonic acid.

Town Gas. Fuel gases containing more than 50% H2 and 10-30% methane together with CO and N2. Obtained mainly by carbonization of coal at 1250°C or by complete gasification of coal

Toxaphene. An insecticide (C10H10Cl8) and a yellow waxy solid. It

Tracer. An additive mainly an isotope of an element used to follow the course of a reaction (or process) with the aid of a counter. E g. radioactive iodine or radioactive oxygen (18O).

Tractor Vapourizing Oil, TVO. A variety of kerosine used for spark engines of low compression ratio.

Trans. A term used in isomerism and represents opposite positions

Transaminase. An enzyme.

Transferin. A protein.

Transormer Oils. They are highly refined oils of low viscosity, high resistance to oxidation used to cooling and insulation in trans-

Transition Elements. A class of elements of the periodic table having partially filled 'd' shell. Three series of elements are: from Sc to Zn, Y to Cd and La to Hg. The first element of series has

1 electron in 'd' shell while the last one has 10 electrons in 'd' shell. These elements have variable valency, coloured ions and tendency to form inorganic complexes. Their salts are paramagnetic.

Transition State (activated complex). The transition state is the high energy configuration through which the reactants must pass before becoming products. The transition state decomposes at a definite rate to yield either the reactants again or the final products. For the reaction,

For the reaction,
$$X+YZ=X...Y...Z \longrightarrow XY+Z$$

Activated complex

The decomposition of transition complex either to reactants or to products depends upon the experimental conditions.

Transition Temperature A temperature at which one (allotropic) form of a substance is converted into another form.

Transmutation. The process of transformation of one element into another by the radioactive decay or by bombardment of the nuclei with particles (e.g., \alpha-particles, neutrons, etc.)

$$\begin{array}{c} 7 \text{ Li} + \frac{1}{1} \text{ p} \longrightarrow 2 \frac{4}{2} \text{ He.} \end{array}$$

Transport Number (t). The fraction of the total current carried by a particular ion in electrolytic conduction.

Tremolite, (OH)2 Ca2 Mg5 (Si4O11)2. An asbestos miner used in acid resistant filters.

Triazines. Compounds containing sym-C₃N₃ rings.

Tribasic Acid. Describing an acid containing three replacable hydrogen atoms in the molecule, e g., H₃PO₄.

1, 1, 1-Trichloroethane, CCl₃.CH₃. A solvent and a metal cleaner prepared by chlorination of ethane.

Frichloroethene, CHCl=CCl₂. A colourless liquid (B.P. 87°C).

Manufactured by the dehydrochlorination of tetrachloro ethane. Used in scouring of wool and as a solvent for

Trichlorofluoromethane, Cl₃CF. A colourless liquid (B.P. 24°C). Used as an integral standard in spectroscopy.

Triclinic System. A crystal system.

Tridymite, SiO₂. An important constituent of silica bricks, stable between 573°C to 1470°C.

Triethylene Glycol, Triglycol. HO CH2.CH2OCH2CH2OCH2CH2OH. Used as a plasticizer and a humidity controller.

Triethyl Phosphate, $(C_2H_5O)_3P=O$. Colourless liquid (B.P. 216°C). Used as a polar reaction medium.

Triethyl Phosphite, (C₂H₅O)₃P. Colourless liquid (B.P. 156°C) used as a deoxygenating desulphurizing agent.

Trifluoroacetic Acid, CF₃COOH. Colourless liquid (B.P. 72.5°C) fumes in air. A strong acid (pk_a=0.3) used as a catalyst in esterification reaction.

Trigol. A trivial name for triethylene glycol.

Trigonal System. The crystal system with a three fold axis as principax axis.

Tri-iodomethane, CHI3. Iodoform.

Trilene. Trichloroethylene when used as an anaesthetic.

2,4,6-Trinitrotoluene, TNT. A yellow crystalline solid obtatined by nitrating methylbenzene or by direct nitration of toluene with a mixture of HNO₃ and H₂SO₄.

$$\begin{array}{c} \text{CH}_3 \\ \\ \hline \\ \text{HNO}_3 + \text{H}_2\text{SO}_4 \\ \\ \hline \\ \text{NO}_3 \\ \\ \end{array} \begin{array}{c} \text{CH}_3 \\ \\ \text{NO}_2 \\ \\ \end{array}$$

Used as a violent and highly exposive (less sensitive to shock and friction).

Triose. A carbohydrate.

Triphenylene, 9, 10-Benzphenanthracene, C₁₈H₁₂. Exhibit fluorescence in solutions Occurs in coal tar (M.P. 199°C).

Triphenylmethane Dyes. A group of dyestuffs derived from triphenyl methane (Ph₃CH) and diphenyl naphthyl methane.

Triphenyl Methyl, Ph₃C. Isolated in the bimolecular form, hexaphenylethane, a colourless solid, M.P. 145-147°C.

Triphenylphospine, (C₆H₅)₃P. Colourless crystals (M.P. 82°C) used as a stabilizing agent for derivatives of some transition metals.

Triple Bond. A covalent bond containing three bonding pairs of electrons between two atoms.

Triple Point. The only point in a one component system at which three phases gas, liquid and solid co exist e.g., the triple point of ice, water and water vapour is 0.0073°C at 4 torr pressure.

Tripolite. A type of diatomite.

Triptycene, C₂₀H₁₄. Colourless solid (M.P. 255°C).

Trithionic Acid, H₂S₃O₆. A polythionic acid.

Tritium, ³H. A radioactive isotope of hydrogen occurring in natural hydrogen (1 in 10⁷ parts). Tritium decays with emission of low-energy β-radiation to form ³He. Used as a tracer and in thermonuclear reactions. Prepared by the deuterium bambardment of some light atoms.

Triton B. A trade name for benzyl trimethyl ammonium hydroxide

(40% solution in methanol). Used as a catalyst.

Trona, Na₂CO₃, NaHCO₃, 2H₂O. Naturally occurring species formed by evaporation of soda lakes.

Tropine, 3-Tropanol, C₈H₁₅NO. A solid, M.P. 63°C.

Tropolones. 2-Hydroxy derivatives of cycloheptatrienone.

Tropylium ion. The positive ion C₃H₇⁺ having a symmetrical sevenmembered ring of carbon atoms.

Truton's Rule. For liquids. an expression

$\frac{M\Delta H v_{\alpha p}}{T}$ = constant

where $\triangle H_{vap}$ is the latent heat of vapourization of liquid, M is the molecular weight and T is the boiling point of liquid (deg K). Normal liquids have a value of about 88 J mol K⁻¹.

Trypsin. A digestive enzyme formed in pancreas (as trypsinogen).

Tryptophah, α-Amino-β-indolylpropionic acid, C₁₁H₁₂N₂O₂. An essential amino acid.

Tung Oil, China-wood Oil. An oil which dries rapidly and obtained from the nuts of Aleurites cordata and Aleurites fordii.

Tungstates. Compounds of W obtained by dissolving WO₃ in NaOH solution followed by cation exchange, eg., Na₂WO₄, 2H₂O. Normal tungstates contain (WO₄)²⁻ lons.

Tungsten, W. At. No. 74, At. wt 183.8, M.P. 3410°C, important ores are: wolframite (Fe, Mn) WO₄, Scheelite (CaWO₄) and Stolzite (PbWO₄). Used to form steel alloys, in electric lamps, in electric contacts, etc. Dissolves in Na₂O₂ or KNO₃—NaOH. Belongs to Group VI and shows oxidation states from +6 to -2.

Tungsten Alloys

Tungsten steels—contain upto 18% W, Steelite—contains W, Cr and Cu and is a very hard alloy, and Cu-W and Ag-W alloys—used for making electric contacts. Tungsten bronzes—MnWO₃ (M_n=unipositive metal 0 < n < 1) Coloured compounds.

- Tungsten Carbides (W2C and WC). Produced by heating W powder with C. Extremely hard and are used for making cutting tools.
- Turbidimetry. A technique of quantitative analysis used to estimate the absorption by a colloidal dispersion of precipitate with the aid of spectrophotometry.
- Turbidity Indicator. An indicator (e g., weak organic acid) used for the titration of glycine (where the change in pH near equiva-
- Turkey Red Oil. A sulphonated castor oil.
- Turpentine. A group of light volatile essential oils obtained by distillation from coniferous trees. B.P. 150°C, D 0.85. Used
- Turquiose, Al₂(OH)₃PO₄, H₂O, plus some Cu, used as a gun stone.
- Tyndall Effect. When a beam of light is led into a colloid, its path becomes visible due to scattering caused by the particles of the disperse phase. This is known as Tynda!! effect. When a convergent beam falls on colloid, a bright blue cone is observed called the Fyndall conc. The effect is greatest for light of short
- Tyrian Purple, C₁₆H₈Br₂N₂O₃. A purple dye.
- Tyrosine, p-Hydroxyphenylalanine, 2-amino-3 (4-hydroxyphenyl) propanoic acid, C₉H₁₁NO₃. M.P. 314-318°C. A least soluble amino acid obtained from proteins.

U. Uranium.

Ubiquinones, Conenzymes Q. A family of 1, 4-benzoquinones which

Ulexite, NaCaB₅O₉, 8H₂O. A source of borate.

- Ulmann Reaction. A reaction for the synthesis of diaryls by the condensation of aromatic halides with themselves or other aromatic halides in the presence of metals, e.g., Cu-powder.
- Ultracentrifuge. A high speed device used for separating out very small particles or to carry out sedimentation of colloidal sols by putting them to large forces. Used to find molecular weights of big molecules, e.g., proteins.
- A technique of filtration in which particles of colloidal dimensions are separated from molecular and ionic sub-Ultrafiltration. stances by drawing the sol liquid through a membrane containing very small capillaries. Also used for sedimentation of large organic molecules in aqueous solution and of heavy inorganic
- Ultramarine. A blue colour aluminosilicate obtained from china clay, S and Na₂CO₃, SiO₂ etc. (a pigment). Contains (Al, Si)O₂ framework together with S or S2.
- Ultramicroscope. A device for examining particles of size 5 to 10 m μ .
- Ultrasonics. A type of vibrations (v≈2×16⁵ cycles per second) produced by applying alternating currents to quartz, tourmaline or rochelle salt or by the effect of oscillating magnetism on a rod of magnetic material immersed in a liquid. Used to investigate fast reactions, for cleaning surfaces, to bleach solutions of coloured dyestuffs, etc.
- Ultra-Violet Absorbers (light stablizers). Compounds which absorb u.v. light and dissipate energy without producing any harm.
- Ultra-Violet Light (u.v.). A term of electromagnetic radiation with wavelength (λ) less than 3600°A. It possesses greater energy than visible radiation. Used to induce photochemical reactions (less penetrating power). Ordinary glasses are nott ransparent to these waves. Quartz is used for making lenses and prisms for
- Umber. A brown powder (pigment) occurring naturally as Fe(OH)2. (containing MnO2).
- Uniaxial. Crystals having one principal axis.
- Unimolecular Films. Unimolecular monolayers obtained by dispersing some insoluble oils and fats (e.g., stearic acid) on the surface
- Unimolecular Reaction. A reaction in which only one reactant (say) molecule) is involved, e.g., a radioactive decay

molecule) is involved, e.g., a rate
$$Rn + \alpha$$

$$Ra \longrightarrow Rn + \alpha$$

$$PCl_5 \longrightarrow PCl_3 + Cl_2$$

Unit Cell. The smallest portion of the space lattice which, when moved in different directions, distances equal to its own dimensions along those directions, develops the whole of the

- Unsaturated Polyesters. Low M.wt. polymers with carboxylic ester grouping and double bonds. Used as co-polymers with monomers in free radical polymerization to form very strong cross-
- Uracil, 2, 6-Dioxytetrahydropyrimidine, C₄H₄N₂O₂. A constituent of ribose nucleic acid (M. P. 280°C), C₄H₄N₂O₂. A constituent of for ribose nucleic acid (M.P. 280°C). Used as a diuretic and for
- Uranium, U. At No. 92, At. wt. 238.029, M.P. 1132.3°C, D 19.07. A toxic radioactive silvery element of actinoid series of metals. Occurs as three isotopes, e.g., 234U (0.005%), 235U (0.0171%) and 288 U (99.283%) in pitchblende U₃O₈), uraninite (UO₂+X), carnotite (KOO₂VO₄, 1.5 H₂₀). Prepared by reduction of UF₄ with Mg at 700°C. Fossil U is used in nuclear weapons. 238 U is a source of fissionable plutonium.
- Uranium Oxides. U forms oxides such as UO (semi-metallic), UO2, U_3O_8 (the stable oxide in air), U_4O_9 , $U_{2\cdot 25}-U_{2\cdot 40}$ and UO_3 .
- Uranyl Derivatives. Compounds of U(VI) containing the (linear) UO2 grouping. Most uranyl derivatives have a strong yellow colour and exhibit bright green fluoresence when irradiated.
- NH. Urea, CH₄N₂O₂, C=O (carbamide). A white crystalline solid,
- (M.P. 132°C) first obtained by Wohler (1828) by evaporating a solution of ammonium cyanate. Manufactured by reacting CO₃ with ammonia at 200°C and 400 at m. pressure. It occurs in urine of mammals and blood of mammals. Used as a fertilizer.
- Urea Adduction. A process of separating normal alkanes from
- Urea Cycle. A cyclic process taking place in liver to excrete excess

Urease. An enzyme.

Ureids. Compounds of urea and organic acids, e.g., barbituric acid.

Urethane, Ethyl Carbamate, NH2COOC2H5. Used as anaesthetic.

Urethanes. Esters of carbonic acid (NH2COOH).

Uric Acid, 2, 6, 8-Trihydroxypurine. A colourless powder. A dibasic Acid, 2, 6, 8-17mys two series of salts. Prepared by fusing

Uricase. An enzyme present in kidney and liver. UV. Ultra violet.

- Trans-11-Octadecenoic Acid, CH2(CH2)5 CH= V. Vanadium CH(CH₂)₉.COOH). An only naturally occurring fatty acid Vaccenic Acid, (M.P. 43-44°C). Present in fats and milk.
- Vacuum. Describing a perfect vacuum with no matter at all.
- Vacuum Pump. A device to extract gas from a space where the pressure is appreciably below the atmospheric pressure and
 - Vacuum Distillation. The distillation of liquids under reduced pressure, so that the boiling point is lowered.
- Valency. A term used to designate the number of bonded neighbour atoms. It is generally equal to either the number of valency electrons or eight minus the number of valency electrons.
- Valency Electrons. The outer electrons in an atom that can determine the chemical reactivity of an element (even geometry of the compound formed by the element).
- Valency, Theory of. Simple valency theory is based on the fact that elements tend to gain, lose or share electrons to attain completion of their outer electron shells. (Noble gas structure).

The electrovalent bond is formed by electrostatic attraction between oppositely charged ions, e.g., NaCl covalent bonds are formed by sharing of electrons between atoms, e.g., H2. Coordinate bonds are formed by the sharing of electrons, both electrons being donated by the same atom, e.g., NH4+ ion.

The shape of covalent compounds are determined by the tendency of bonding pairs to be as far apart as possible whereas lone pairs have a greater effect that bonding (VSEPR).

Valeric Acids, Pentanoic Acids, C₅H₁₀O₂. n-Valeric acid, CH₂CH₂. CH₂COOH. A colourless liquid (B.P. 186°C) prepared by oxidizing n-amyl alcohol or from Grignards' reagent plus n-butyl iodide.

- Iso-Valeric Acid, (CH₃)₂CH.CH₂COOH. A colourless liquid (B.I 177°C). Prepared by oxidizing isoamyl alcohol.
- Valine, 2-Amino-3-Methylbutanoic Acid, C₅H₁₁NO₂. An amino acid obtained by the hydrolysis of proteins. M.P. 315°C.
- Valium. Diazepam (a sedative).
- Vanadium, V. At. No. 23, At. wt. 50.94, M.P. 1890°C, D 5.96. A transition element of Group V occurring naturally as patronite (V₂S₅+5), carnotite (KUO₂VO₄, 1.5H₂O) and vanadinite [Pb₅. (VO₄)₃Cl]. Electronic configuration ²d³ 4s², shows oxidation states from states from -1 to +5 (+5 in covalent or in complexes). Forms coloured ions, eg., V (IV) forms blue (in aqueous solutions) V (III) 2000 (ig., V (IV) forms blue (in aqueous law-red solutions). V (III) green (in aqueous solution), V (V) yellow-red in oxides. Used as a steel oddition in oxides. Used as a steel additive and in V-Al alloys.
- Vanadian (V) Oxide, V₂O₅. (Vanadium pentaoxide). Orange-yellow oxide (heat NH₄VO₃). Dissolves in alkalis to form vandates (v)-Used as a catalyst (Control of the control of the co Used as a catalyst (Contact process).
- Vanadyl Species. Compound having the blue VO2+ group, e.g.,
- Van Arkel-de Boer Process. A method for purification of metals.
- waal's forces between the coloridate adsorption having vander waal's forces between the adsorbate and adsorbent.
- van der Waal's Equation. A modified form of state for gases:

$$\left(P+n^2\frac{a}{\sqrt{2}}\right)(V-nb)=nRT$$

- where a and b are van der Waals constant of the gas (specific constants) and n is the constants) and n is the number of moles of gas (b is the covolume'—equal to 4 times the number of moles of gas (b is the 'covolume'—equal to 4 times the actual volume of molecules and $\frac{a}{\nabla^2}$ is the pressure correction).
- van der Waal's Forces. The weak intermolecular forces between the molecules which arise from weak electronic coupling.
- Vanillin, 4-Hydroxy-3-Methoxybenzaldehyde, C₈H₈O₃. Fine white needles (M.P. 82°C) of strong vanilla odour. Occurs in nature. Manufactured from the lignosulphonic acid (a by-product in wood pulp manufacture). Used as a flavouring agent and for
- Van't Hoff Factor, (i). The ratio of the number of particles present in as olution to the number of undissociated molecules added. Used in the study of colligative properties of solutions. For example, vant Hoff factor i is the ratio of observed osmotic pressure and the theoretically calculated or ideal osmotic pressure.

van't Hoff Isochore. The equation

$$\frac{\mathrm{d} \ln K}{\mathrm{d} T} = \frac{\triangle H}{RT^2}$$

where K is the equilibrium constant of a reversible reaction, AH the enthply of reaction, T the absolute temperature and R the gas constant, Used to calculate enthalpies of reactions.

van't Hoff Isotherm. For a reversible chemical reaction such as aA+B = cC+dD

the free energy change, $\triangle G = -RT \ln K + RT \frac{[C]^c \cdot [D]^d}{[A]^a B^b}$ where

R is gas constant, T the absolute temperature. K the equilibrium contant and [A][B], etc., the activities of A, B, etc., at equilibrium conditions.

Vapourization. A process by which a solid or liquid is converted into a gas or vapour by heating.

Vapour. A gas formed by the vapourization of a solid or liquid.

Vapour Density. The pressure exerted by liquid or solid at equilibrium and at specifie I temperature.

Vaseline (soft paraffin). Yellow or white semisolid containing mixtures of hydrocarbons of the paraffin series (C15H32 to C₂₀H₄₂). Prepared from high-boiling fractions of petroleum or shale oil. Used as a base for ointment and as dressing for wounds.

Vaska's Compound. A trans-[fr (CO) (Ph₃P)₂Cl] compound.

Vat Dyes. A class of water insoluble dyestuffs applied to fibres by first reducing them to derivatives that are soluble in dilute alkalis On reoxidation (with air), the insoluble dyestuffs is precipitated within the fibre.

Velocity of Reaction. The amount of reactants transformed or of products formed, per unit time.

Veratraldehyde, C9H10O3 Obtained by methylation of vanillin which on further oxidation forms veratric acid

Verbenone, C₁₀H₁₄O. A cyclic ketone.

Verdet's Constant. A constant used in magnetic polarization of light.

Verdigris. A green product formed on Cu or bronze when exposed to air (basic copper carbonate).

Veridian, CraO3.

Vermiculite A hydrated silica clay.

Vermillion Red, HgS. A red pigment which occurs in nature as cinnabar.

Vibrational Spectrum. See spectrum.

- Vic. A prefix indicating that in an organic compound, the substituent atoms or groups are bounded to adjacent C-atoms.
- Victor Meyer Method. A method for determining vapour densities (or M. wt.) in which a given weight of sample is vapourized and the volume of the air displaced is measured.
- Vinegar. A dilute (4-10%) solution of ethanoic acid (CH₃COOH) obtained by the oxidation of alcoholic liquors by Acetobacter species such as Acetobacter aceti.
- Vinyl Ethyl. The group CH₂=CH-.
- Vinyl Acetate, Vinyl Ethanoate, CH₂=CHOOC.CH₃. Colourless . liquid (B.P. 75°C). Manufactured by vapour phase reaction of ethyne and ethanoic acid (Zinc ethanoate catalyst) at 200°C. Used in polymer manufacture.
- Vinylation. An addition reaction across the triple bond (e.g., ethyne and carboxylic acids, amines etc.,) to form vinyl compounds, e.g., H-C≡C-H+HX → CH₂=CHX.
- Vinyl Chloride Polymers. Polymers, having (CH₂—CHCl)_n units including potyvinyl chloride, PVC etc. Used in rigid materials, flooring, electrical coverings, packaging, clothing etc.
- Vinyl Ester Polymer. Polyvinyl acetate is the important polymer of etc.

 this class. Used in adhesives, as thickness, solvents, plasticizers
- Vinyl Ethers, ROCH= CH_2 . A group of ethers prepared by reacting alkynes (C_2H_2) and alcohols at about 150°C under pressure in MeOCH= CH_2 .
- Vinylidene Chloride Polymers, Saran Polymers. Polymers produced by the free radical polymerization of CH₂=CCl₂. Have high thermal stability and low permeability to gases.
- Violaxanthin, C₄₀H₅₆O₄. A carotenoid pigment of reddish brown colour (M.P. 20°C).
- Virial Equation. An equation for a gas

$$\frac{PV}{RT} = 1 + \frac{B}{V} + \frac{C}{V^2} + \dots$$

where B and C are second and third virial co-efficient respec-

- Viscosity. A property of fluids. All fluids show a resistance to flow. Viscosity measurements are used to find the molecular weights
- Viscosity, Coefficient of (η) . It is the tangential force required per unit area to maintain unit difference of difference of velocity between two layers unit distance apart. It is expressed in units of poise (dynes-sec per cm).

Vitamins. Substances which are essential constituents of food of animal. They do not include proteins, carbohydrates, fats and mineral salts. They play an important role in animal metabolic process.

Vitamio A. A fat soluble vitamin present in animal fats, butter, yolk of egg, in fish-liver oils. It has empirical formula $C_{90}H_{30}O$. Carotene is converted to vitamin A in liver. Two molecules of vitamin A are formed from one molecule of β-carotene. Pale yellow needles (M P. 64°C).

Vitamin B.

Vitamin B₁ - thiamine,

Vitamin B₃ - riboflavin,

Vitamin B₁₂ - Cyanocobalamine,

Vitamin Bo - folic acid,

Vitamin B₆ - Pyridoxine.

Vitamin B₁₂, Cyanocobalamine, C₆₃H₉₀CoN₄O₁₄P. Dark red crystals. Occurs in liver. Produced by the growth of some microorganisms. Its deficiency is due to failure to absorb B18 from stomach.

Vitamin C. Ascorbic acid.

Vitamin D. A fat soluble vitamin present in animal fats, milk, butter and eggs Its deficiency lead to the development of rickets.

Vitamin E. An essential vitamin for fertility and reproduction. Present in seed embroys and their oils, and green leaves.

Vitamin K. A fat soluble vitamin present in lever fats, vegetables and in cereals. Deficiency leads to haemorrhages due to a failure of blood to clot properly.

Vitellin. Chief protein present in egg yolk.

Volhard Method. A volumetric method for the estimation of Ag using NCS- ions in the presence of Fe3+ (deep red colour at the end point).

Vulcanization. A process of improving the quality of rubber (hardness and resistance to temperature changes) by heating it with S (at 150°C) in the presence of accelerators (e.g., Se, Te etc.).



W. Tungsten.

Wacker Process. A method of preparing ethanal from ethane,

$$C_2H_4+O_3 \xrightarrow{Cu^2/Pd^{2+}} CH_3CHO$$

- Walden Inversion. A reaction in which an optically active compound figuration has been inverted. For example S_N2 reactions. It acid from (+) malic acid.
- Warfarin, C₁₉H₁₆O₄. An anti-coagulant rodenticide. Its derivatives
- Washing Soda, Na₂CO₃, 10H₂O. Sodium carbonate.
- Water, H₂O. M.P. 0°C, B.P. 100°C, D. at 0°C 0.99987. A hydride of oxygen which exists as single H₂O in the gaseous state. With a residual order. The atoms do not lie in single plane, atom is 105°, the distance between atoms and central oxygen oxygen atom is 0.99 mm. It is polar in nature. Behaves as a Forms hydrates with many salts such as CuSO₄, 5H₂O. Forms hydroxides with electropositive metals, e.g., Na, K, etc. Used as a powerful solvent (high dielectric constant).
- Water Gas. A mixture of CO and H₂ produced when steam is passed over red hot coke.
- Water of Crystallization. Water present in definite proportions in crystalline compounds, e.g., Na₂CO₃. 10H₂O. It can be removed
- Water-Proffing. A process by which a fabric is made impermeable to water by impregnation with oils, varnishes, rubber, etc.
- Wavelength, (λ) . The distance between corresponding points in the profile of wave motion.

- Wave Mechanics. A system of mechanics that is developed from quantum theory and is used to explain the behaviour of atoms, molecules, etc. It is based on de Broglie's idea that particles can have wave like properties. Wave mechanics is a branch of quantum mechanics and is the basis of modern theory.
- Wave Number. The number of waves in a given length (generally 1 cm). A reciprocal of wavelength.
- Waxes. Any organic material having wax like properties. Used in paper coating polishes, insulators, leathers, etc.
- Weber, Wh. A unit of magnetic flux, equal to the magnetic flux that linking a circuit of one turn, produces an em.f. of one volt when reduced to zero at a uniform rate in one second.

1Wb=1Vs

- Weed Killers. Herbicides used to control selectivily unwanted vegetation on agricultural land.
- Weston Cadmium Cell. A standard cell Cd (Hg)/CdSO₄, ⁵/₃H₂O(s), || Hg₂SO₄(s)/Hg. The electrolute is a saturated solution of CdSO4.

E20°c=1.01485 volts

- Wet-Bulb Thermometer. A thermometer for measuring wet-bulb temperature in which bulb is surrounded by wicks (to keep damp), and the liquid is allowed to evaporate adiabatically.
- Wetting Agents. Additives which wet the surfaces because of its powerful intermolecular attractive forces, e.g., water.

White Lead, Pb3(OH)2(CO3)2.

White Oils. Refined lubricant oils used in toilet preparations.

White Spirits. Mineral Solvents. A mixture of hydrocarbons (B.P. 130-220°C). Used as paint thinners and as a solvent.

Whiting. A variety of CaCO₃.

Wilkinson's Catalyst. A catalyst used for homogeneous hydrogenation. [Rh(PhaP)aCl].

Williamson's Synthesis. A method for the preparation of mixed ethers by refluxing a haloalkane and an alcoholic solution of sodium alkoxide

R'Cl+RO-Na+ → R'OR+NaCl

(A nucleophilic substitution).

Witherite, BaCO₃. A mineral (white) of barium carbonate.

Wittig Reaction. A reaction between an alkylidene phosphorane and an aldehyde or ketone to form an alkene.

WLN. Wiswesser line notation.

Woframite, (Fe, Mn) WO4.

Wood. A complex substance with cell walls having carbohydrates (cellulose) and lignin plus some inorganic and organic materials. Used for constructing furniture.

Wood's Metal. A fusible alloy containing Bi (50%), Pb (25%), Sn (12.5%) and Cd. Used in fire protection devices (M.P.70°C).

Wool. A natural protein fibre (keratin).

Work Function. The energy needed to remove an electron from the highest occupied level inside a solid to a point in vacuo outside

Wrought Iron. A low carbon steel. Used to make girders.

Wurtzite. A variety of ZnS.

Wurtz Synthesis. A reaction for preparing alkanes by refluxing (RX) with Na in dry ether

2RX+2Na → RR+2NaX

- Xanthates. The salts or esters of unstable acids, ROC(S)SH, where R may be either an alkyl or any group. Used in curing and vulcanizing rubber. They may be coloured compounds.
- Xanthine, 2, 6—Oxypurine. C₅H₄N₄O₂. A colourless powder which
- Kanthone, C₁₃H₈O₂. Colourless crystals (M.P. 174°C). Obtained by heating phenyl salicylate. A parent substance of xanthone
- Xanthophyll, C40H86O2. Yellow pigments found with carotene in most green plant material.

Xe. Xenon.

Xenon, Xe. At. No. 54, At. wt. 131.30, M.P. -111.9°C, B.P. -107.1°C A noble gas with electronic configuration 55°p6. Used in lamps, discharge tubes and in bubble chamber.

Xenvlamine. C12H11N.

Xerogels. A technique for classification of gels.

Xerography. A process in which a photo-conductor on a surface is charged, exposed to the image which discharges all except the image, removal of photo-conductor from the non-image areas, developing the image with an oppositively charged pigment, transferring the image to paper electrostatically and fixing the image on paper by baking.

X-ray. An electromagnetic radiation of wavelength 10-8 cm (1A°). Generated by bombarding solids wish electrons. Each element has a characteristic X-rays spectrum. Used in crystallography

X-ray Spectroscopy. Analytical technique involving the irradiation of sample with X-rays.

X-ray Tube. An apparatus used in X-ray production.

Xylene, Dimethylbenzene, C₈H₁₀. A colourless liquid and a mixture of three isomers, e.g., o-Xylene (B.P. 144°C), m-Xylene (B.P. 139°C) and p-Xylene (B.P. 138°C). Obtained by reforming naphthalene in presence of H2. Also prepared from coal-tar.

Xylic Acid. A name for 2, 4-dimethylbenzoic acid.

D-Xylose, Wood Sugar, C₅H₁₀O₅. A pentose sugar (M.P. 144°C).



Y. Yttrium.

Y Alloy. An alumium alloy.

Yellow Ammonium Sulphide. A poly ammonium sulphide.

Ylides. An internal salt, e.g.,

 $R_1C=PPh_3 \leftrightarrow R_2C^{-2}P^+Ph_3$.

Ytterbium, Yb. At. No. 70, At. wt. 173.04, M P. 824°C, D. 6.98. A lanthanide used to improve mechanical power of steel and as synthetic gem stone. Shows +2 and +3 oxidation states.

338 Yttrium Y

Yttrium, Y. At. No. 39, At. wt. 88.91. M.P. 1523°C, D. 4.472. An element of Group III with electronic configuration 5S' 4d1. Important ore is gadolinite. Y-Fe garnets are used as micro-

Zein. A protein present in maize. Used in the plastics, paints etc.

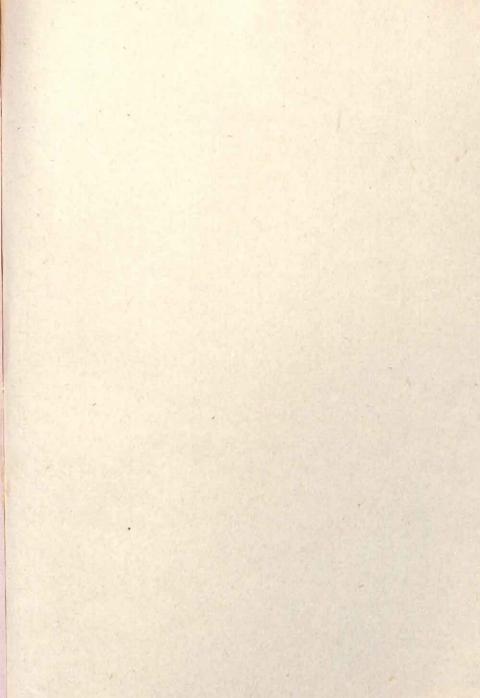
Zeisel Reaction. A reaction of ether with excess conc. HI.

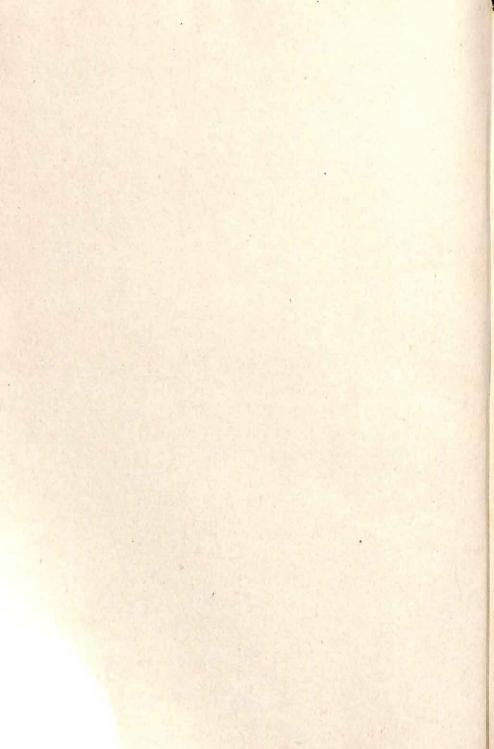
 $ROR'+2HI \longrightarrow RI+RI'+H_2O$

Zeise's Salt, K [Pt Cl₃ (C₂H₄)].

- Zeolites Aluminosilicates which occur in nature and contain (Si, Al)_nO_{2n} frameworks with a negative charge netralized by cations present in the cavities. Used as a water softner and
- Ziegler Natta Polymerisation. A process of polymerisation of olefins
- Zinc, Zn. At. No. 30, At. wt. 65.38, M.P. 419.58, D. 7.13. A transition metal of Group II with electronic configuration 3d¹⁰4s³. Occurs as sphalerite (Zn, Fe) S, smithsonite (Zn CO₃) Franklite (Zn Fe₂ O₄). Dissolves in dilute acids and hot solutions of alkalis. Used in anodes for protection against
- Zinc Am. Igam. A solid Zn—Hg solution (a reducing agent).
- Zincates. Ions such as [Zn (OH)₄]⁻² formed by the action of excess
- Zinc Blende, ZnS. The Zn and S are tetrahedrally co-ordinated.
- Zinc Borates. Compounds of indefinite composition. Used for fire
- Zinc Chloride, ZnCl₂. White solid (Zn+HCl). Used as wood
- Zinc Chromate. A yellow pigment (ZnSO₄+K₂Cr₂O₇) used in rust

- Zinc Dithionite, ZnS₂O₄. (SO₂+aqueous Zn dust).
- Zinc Fluoride, ZnF₂. 4H₂O. Forms ZnF₃ at 100°C (Zn+F₃ or HF). Used as a wood preservative.
- Zinc Hydroxide, Zn (OH)2. A rubber filler and an adsorbent in
- Zinc Nitrate, Zn (NO₃)₂, xH₂O.
- Zinc Oxide, ZnO. A powder white (cold) yellow (hot). Obtained by burning Zn in O₃. Used as a disinfectant in pharmacy.
- Zinc Sulphate, White Vitriol, Zn SO4. Forms, 7, 6, 1 and 0 hydrates (ZnO+H₂SO₄). Used in textiles.
- Zingiberene, C₁₅H₂₄. A constituent of ginger oil (B.P. 134°C/14 nm).
- Zircon, Zr Si O4. A mineral of Zr.
- Zirconium, Zr. At. No. 40, At. wt. 91.22, 1852°C, D. 6.506. A transition element of Group IV, electronic configuration $5s^24d^3$. Occurs as baddelyite (ZrO₂) and zircon (ZrSiO₄). Resistant to corrosion but dissolved by aquaregia. Forms zirconium compounds with +4 oxidation state. Its alloys are used in reactor construction, and in super conducting magnets (Nb-Zr alloy). Zn. Zinc.
- Zone Refining. A process of refining metals and some organic and inorganic compounds depending on the difference in solubility of impurities in the liquid and solid states. Used in
- Zwitterion (ampholyte ion). An ion that has both positive and negative charge. An electrically neutral ion, e.g., amino acid glycine exists at isoelectric ion in solution as the Zwitter ion
- Zymase. An enzyme present in yeast. Used to convert glucose to







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